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(54) Title: **METHODS AND COMPOSITIONS FOR MODULATING INTERLEUKIN-21 RECEPTOR ACTIVITY**

(57) Abstract: Methods and compositions for modulating interleukin-21 (IL-21)/IL-21 receptor (MU-1) activity using agonists or antagonists of IL-21 or IL-21 receptor ("IL-21R" or "MU-1"), are disclosed. IL-21/IL-21R antagonists can be used to induce immune suppression *in vivo*, e.g., for treating or preventing immune cell-associated pathologies (e.g., pathologies associated with aberrant activity of one or more of mature T cells (mature CD8+, mature CD4+ T cells), mature NK cells, B cells, macrophages and megakaryocytes, including transplant rejection and autoimmune disorders). IL-21/IL-21R agonists can be used by themselves or in combination with an antigen, e.g., as an adjuvant (e.g., a vaccine adjuvant), to up-regulate an immune response *in vivo*, e.g., for example, for use in treating cancer and infectious disorders.

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METHODS AND COMPOSITIONS FOR MODULATING INTERLEUKIN-21 RECEPTOR ACTIVITY

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FIELD OF THE INVENTION

The present invention relates to methods and compositions for modulating interleukin-21 (IL-21)/IL-21 receptor (MU-1) activity using IL-21 receptor agonists and antagonists. The methods and compositions disclosed herein are useful as immunotherapeutical agents.

BACKGROUND OF THE INVENTION

Human IL-21 is cytokine about a 131-amino acids in length that shows sequence homology to IL-2, IL-4 and IL-15 (Parrish-Novak *et al.* (2000) *Nature* 408:57-63).
Despite low sequence homology among interleukin cytokines, cytokines share a common fold into a "four-helix-bundle" structure that is representative of the family. Most cytokines bind either the class I or the class II cytokine receptors. Class II cytokine receptors include the receptors for IL-10 and the interferons, whereas class I cytokine receptors include the receptors for IL-2-IL-7, IL-9, IL-11-13, and IL-15, as well as hematopoietic growth factors, leptin and growth hormone (Cosman, D. (1993) *Cytokine* 5:95-106).

Human IL-21R is a class I cytokine receptor that is expressed in lymphoid tissues, in particular by NK, B and T cells (Parrish-Novak *et al.* (2000) *supra*). The nucleotide and amino acid sequences encoding human interleukin-21 (IL-21) and its receptor (IL-21R) are described in WO 00/53761; WO 01/85792; Parrish-Novak *et al.* (2000) *supra*; Ozaki *et al.* (2000) *Proc. Natl. Acad. Sci. USA* 97:11439-11444. IL-21R has the highest sequence homology to IL-2 receptor β chain and IL-4 receptor α chain (Ozaki *et al.* (2000) *supra*). Upon ligand binding, IL-21R associates with the common-gamma cytokine receptor chain (γ_c) that is shared by receptors for IL-2, IL-3, IL-4, IL-7, IL-9, IL-13 and IL-15 (Ozaki *et al.* (2000) *supra*; Asao *et al.* (2001) *J. Immunol.* 167:1-5). The widespread lymphoid distribution of IL-21R suggests that IL-21 may play a role in immune regulation. Indeed, *in vitro* studies have shown that IL-21 significantly modulates the function of B cells, CD4⁺ and CD8⁺ T cells, and NK cells (Parrish-Novak

et al. (2000) *supra*; Kasaian, M.T. *et al.* (2002) *Immunity*. 16:559-569). Nevertheless, evidence supporting a regulatory effect of IL-21 *in vivo* is limited.

SUMMARY OF THE INVENTION

Methods and compositions for modulating the activity of, and/or an interaction
5 between, an interleukin-21 (IL-21) and an IL-21 receptor (also referred to herein as "IL-21R" or "MU-1") using agonists or antagonists of IL-21 or IL-21R are disclosed (also referred to herein as an "IL-21/IL-21R agonist" or "agonist," and "IL-21/IL-21R antagonist" or "antagonist," respectively).

In one embodiment, Applicants have shown that reducing IL-21R activity by
10 using an IL-21 antagonist, e.g., a fusion protein that includes the extracellular domain of the IL-21R fused to an Fc immunoglobulin region, ameliorates inflammatory symptoms in collagen-induced arthritis (CIA) animal models (Example 7). Expression of IL-21R mRNA is upregulated in the paws of CIA mice (Example 8). Accordingly, antagonists of IL-21/IL-21R activity can be used to induce immune suppression *in vivo*, e.g., for treating
15 or preventing immune cell-associated pathologies (e.g., pathologies associated with aberrant activity of one or more of mature T cells (mature CD8+, mature CD4+ T cells), mature NK cells, B cells, macrophages and megakaryocytes, including transplant rejection and autoimmune disorders, e.g., arthritis (including rheumatoid arthritis).

Accordingly, in one aspect, the invention features a method of treating (e.g.,
20 curing, suppressing, ameliorating, delaying or preventing the onset of, or preventing recurrence or relapse of) or preventing an immune cell-associated disease, e.g., rheumatoid arthritis, in a subject, the method includes: administering to the subject an IL-21/IL-21R antagonist, in an amount sufficient to inhibit or reduce immune cell activity and/or cell number, thereby treating or preventing the immune cell-associated disease,
25 e.g., rheumatoid arthritis.

The IL-21/IL-21R antagonist can be administered to the subject, alone or in combination, with other therapeutic modalities as described herein. Preferably, the subject is a mammal, e.g., a human suffering from an immune cell-associated pathology (e.g., pathology associated with aberrant activity of one or more of mature T cells
30 (mature CD8+, mature CD4+ T cells), mature NK cells, B cells, macrophages and megakaryocytes, including transplant rejection and autoimmune disorders. For example, the method can be used to treat or prevent, in a subject, an immune cell-associated

disorders, e.g., a disorder chosen from one or more of: transplant rejection or an autoimmune disorder (e.g., including, for example, diabetes mellitus (type I), arthritis (including rheumatoid arthritis, juvenile rheumatoid arthritis, osteoarthritis, psoriatic arthritis, and ankylosing spondylitis), multiple sclerosis, myasthenia gravis, vasculitis, systemic lupus erythematosus, autoimmune thyroiditis, dermatitis (including atopic dermatitis and eczematous dermatitis), psoriasis, scleroderma, asthma, allergy, IBD or Crohn's disease). Treatment of an arthritic disorder, e.g., a disorder chosen from one or more of rheumatoid arthritis, juvenile rheumatoid arthritis, osteoarthritis, psoriatic arthritis, or ankylosing spondylitis (preferably, rheumatoid arthritis) using the IL-21- or IL-21R antagonists of the present invention is preferred.

In one embodiment, the IL-21/IL-21R antagonist interacts with, e.g., binds to, IL-21 or IL-21R, preferably, mammalian, e.g., human IL-21 or IL-21R (referred to herein as an "IL-21 antagonist" and "IL-21R antagonist," respectively), and reduces or inhibits one or more IL-21 and/or IL-21R activities. Preferred antagonists bind to IL-21 or IL-21R with high affinity, e.g., with an affinity constant of at least about 10^7 M^{-1} , preferably about 10^8 M^{-1} , and more preferably, about 10^9 M^{-1} to 10^{10} M^{-1} or stronger.

For example, an IL-21/IL-21R antagonist can reduce and/or inhibit IL-21R activity by neutralizing IL-21. In one embodiment, the antagonist can be a fusion protein that includes a fragment of an IL-21R fused to a non-IL21R fragment, e.g., an immunoglobulin Fc region. In other embodiments, the antagonist is an anti-IL21R or anti-IL21 antibody or an antigen-binding fragment thereof, a soluble form of the IL-21 receptor, a peptide or a small molecule inhibitor.

In one embodiment, the IL-21/IL-21R antagonist is an anti-IL21R or anti-IL21 antibody, or an antigen-binding fragment thereof. E.g., the antibody is a monoclonal or single specificity antibody, that binds to IL-21, e.g., human IL-21, or an IL-21 receptor, e.g., human IL-21 receptor polypeptide, or an antigen-binding fragment thereof (e.g., an Fab, F(ab')₂, Fv or a single chain Fv fragment). Preferably, the antibody is a human, humanized, chimeric, or *in vitro* generated antibody to human IL-21 or human IL-21 receptor polypeptide. Preferably, the antibody is a neutralizing antibody.

In other embodiments, the IL-21/IL-21R antagonist includes full length, or a fragment of an IL-21 polypeptide, e.g., an IL-21 receptor-binding domain of an IL-21 polypeptide, e.g., a human IL-21 polypeptide (e.g., a human IL-21 polypeptide as

described herein having an amino acid sequence shown as SEQ ID NO:19) or a sequence at least 85%, 90%, 95%, 98% or more identical thereto; or encoded by a corresponding nucleotide sequence shown as SEQ ID NO:18 or a sequence at least 85%, 90%, 95%, 98% or more identical thereto. Alternatively, the antagonist includes full length (e.g.,
5 from about amino acids 1-538 or 20-538 of SEQ ID NO:2; or from about amino acids 1-529 or 20-529 of SEQ ID NO:10), or a fragment of an IL-21 receptor polypeptide, e.g., an IL-21-binding domain of an IL-21 receptor polypeptide, e.g., a soluble fragment of an IL-21R (e.g., a fragment of an IL-21R comprising the extracellular domain of murine or human IL-21R; e.g., from about amino acids 1-235, 1-236, 20-235, 20-236 of SEQ ID
10 NO:2 (human), or from about amino acids 1-236, 20-236 of SEQ ID NO:10 (murine), or encoded by the corresponding nucleotides of SEQ ID NO:1 or 9, or a sequence at least 85%, 90%, 95%, 98% or more identical thereto.

In one embodiment, the antagonist is a fusion protein comprising the aforesaid IL-21 or IL-21 receptor polypeptides or fragments thereof and, e.g., fused to, a second
15 moiety, e.g., a polypeptide (e.g., an immunoglobulin chain, a GST, Lex-A or MBP polypeptide sequence). In a preferred embodiment, the fusion protein includes at least a fragment of an IL-21R polypeptide, which is capable of binding IL-21, e.g., a soluble fragment of an IL-21R (e.g., a fragment of an IL-21R comprising the extracellular domain of murine or human IL-21R; e.g., from about amino acids 1-235, 1-236, 20-
20 235, 20-236 of SEQ ID NO:2 (human), or from about amino acids 1-236, 20-236 of SEQ ID NO:10 (murine), or encoded by the corresponding nucleotides of SEQ ID NO:1 or 9, or a sequence at least 85%, 90%, 95%, 98% or more identical thereto) and, e.g., fused to, a second moiety, e.g., a polypeptide (e.g., an immunoglobulin chain, an Fc fragment, a heavy chain constant regions of the various isotypes, including: IgG1, IgG2, IgG3, IgG4,
25 IgM, IgA1, IgA2, IgD, and IgE). For example, the fusion protein can include the extracellular domain of human IL-21R, e.g., from about amino acids 1-235, 1-236, 20-235, 20-236 of SEQ ID NO:2, and, e.g., fused to, a human immunoglobulin Fc chain (e.g., human IgG, e.g., human IgG1 or a mutated form of human IgG1). In one embodiment, the human Fc sequence has been mutated at one or more amino acids, e.g.,
30 mutated at residues 254 and 257 of SEQ ID NO:28, from the wild type sequence to reduce Fc receptor binding. In other embodiments, the fusion protein can include the extracellular domain of murine IL-21R, e.g., from about amino acids 1-236, 20-235 of

SEQ ID NO:10 (murine), and, e.g., fused to, a murine immunoglobulin Fc chain (e.g., murine IgG, e.g., murine IgG2a or a mutated form of murine IgG2a).

The fusion proteins may additionally include a linker sequence joining the first moiety, e.g., an IL-21R fragment, to the second moiety, e.g., the immunoglobulin fragment. In other embodiments, additional amino acid sequences can be added to the N- or C-terminus of the fusion protein to facilitate expression, steric flexibility, detection and/or isolation or purification.

Examples of antagonistic fusion proteins that can be used in the methods of the invention are shown in Figures 7-15. In one embodiment, the fusion protein includes an amino acid sequence chosen from, e.g., SEQ ID NO:23, SEQ ID NO:25, SEQ ID NO:27, SEQ ID NO:29, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:35, SEQ ID NO:37, or SEQ ID NO:39, or a sequence at least 85%, 90%, 95%, 98% or more identical thereto. In other embodiments, the fusion protein includes an amino acid sequence encoded by a nucleotide sequence chosen from, e.g., SEQ ID NO:22, SEQ ID NO:24, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:30, SEQ ID NO:32, SEQ ID NO:34, SEQ ID NO:36, or SEQ ID NO:38, or a sequence at least 85%, 90%, 95%, 98% or more identical thereto. Preferred fusion proteins have the amino acid sequence shown as SEQ ID NO:25 or SEQ ID NO:29 (Figures 8A-8C and 10A-10C, respectively), or a sequence at least 85%, 90%, 95%, 98% or more identical thereto. In other embodiments, the fusion protein includes an amino acid sequence encoded by a nucleotide sequence chosen from, e.g., SEQ ID NO:24 or SEQ ID NO:28 (Figures 8A-8C and 10A-10C, respectively), or a sequence at least 85%, 90%, 95%, 98% or more identical thereto. Most preferably, the fusion protein has the amino acid sequence shown as SEQ ID NO:29, or has an amino acid sequence encoded by a nucleotide sequence shown as SEQ ID NO:28 (Figure 10A-10C).

The IL-21/IL-21R antagonists described herein, e.g., the fusion protein described herein, can be derivatized or linked to another functional molecule, e.g., another peptide or protein (e.g., an Fab' fragment). For example, the fusion protein or an antibody, or antigen-binding portion, can be functionally linked (e.g., by chemical coupling, genetic fusion, non-covalent association or otherwise) to one or more other molecular entities, such as an antibody (e.g., a bispecific or a multispecific antibody), toxins, radioisotopes, cytotoxic or cytostatic agents, among others.

In one embodiment, the IL-21/IL-21R antagonists described herein, e.g., the pharmaceutical compositions thereof, are administered in combination therapy, i.e., combined with other agents, e.g., therapeutic agents, which are useful for treating immune cell-associated pathological disorders, e.g., a disorder chosen from one or more of: arthritis (including rheumatoid arthritis, juvenile rheumatoid arthritis, osteoarthritis, psoriatic arthritis, or ankylosing spondylitis), or scleroderma, systemic lupus erythematosus, or vasculitis; preferably, rheumatoid arthritis). For example, the combination therapy can include one or more IL-21/IL-21R antagonists, e.g., an anti-IL-21- or anti-IL-21R antibody or an antigen-binding fragment thereof; an IL-21R fusion protein; a soluble IL-21R receptor; a peptide inhibitor or a small molecule inhibitor) co-formulated with, and/or co-administered with, one or more additional therapeutic agents, e.g., one or more cytokine and growth factor inhibitors, immunosuppressants, anti-inflammatory agents, metabolic inhibitors, enzyme inhibitors, and/or cytotoxic or cytostatic agents, as described in more herein.

Examples of preferred additional therapeutic agents that can be co-administered and/or co-formulated with one or more IL-21/IL-21R antagonists, include, but are not limited to, one or more of: TNF antagonists (e.g., chimeric, humanized, human or *in vitro* generated antibodies, or antigen-binding fragments thereof, that bind to TNF; soluble fragments of a TNF receptor, e.g., p55 or p75 human TNF receptor or derivatives thereof, e.g., 75 kDTNFR-IgG (75 kD TNF receptor-IgG fusion protein, EnbrelTM), p55 kD TNF receptor-IgG fusion protein; TNF enzyme antagonists, e.g., TNF α converting enzyme (TACE) inhibitors); antagonists of IL-12, IL-15, IL-17, IL-18, IL-22; T cell and B cell depleting agents (e.g., anti-CD4 or anti-CD22 antibodies); small molecule inhibitors, e.g., methotrexate and leflunomide; sirolimus (rapamycin) and analogs thereof, e.g., CCI-779; Cox-2 and cPLA2 inhibitors; NSAIDs; p38 inhibitors, TPL-2, Mk-2 and NF κ b inhibitors; RAGE or soluble RAGE; P-selectin or PSGL-1 inhibitors (e.g., small molecule inhibitors, antibodies thereto, e.g., antibodies to P-selectin); estrogen receptor beta (ERB) agonists or ERB-NF κ b antagonists. Most preferred additional therapeutic agents that can be co-administered and/or co-formulated with one or more IL-21/IL-21R antagonists include one or more of: a soluble fragment of a TNF receptor, e.g., p55 or p75 human TNF receptor or derivatives thereof, e.g., 75 kDTNFR-

IgG (75 kD TNF receptor-IgG fusion protein, EnbrelTM); methotrexate, leflunomide, or a sirolimus (rapamycin) or an analog thereof, e.g., CCI-779.

Applicants have further shown that an IL-21/IL-21R agonist, e.g., an IL-21 polypeptide, stimulates immunity *in vivo* against immunogenic and non-immunogenic tumor cells (Example 9). The increased immunity is due in part by the IL-21-mediated potentiation of mature CD8+ T cell effector function. IL-21/IL-21R agonists can be used by themselves or in combination with an antigen, e.g., as an adjuvant (e.g., a vaccine adjuvant), to up-regulate an immune response *in vivo*, e.g., for example, for use in treating cancer or an infectious disorder in a subject.

Accordingly, the invention provides a method of treating (e.g., curing, suppressing, ameliorating, delaying or preventing the onset of, or preventing recurrence or relapse of) or preventing a cancer or an infectious disorder, in a subject, the method includes: administering to the subject an IL-21/IL-21R agonist, e.g., an agent that increases or potentiates IL-21/IL-21R activity, in an amount sufficient to increase immune cell (e.g., CD8+ cell) activity (e.g., effector cell activity) and/or cell number, thereby treating or preventing said disorder. Exemplary cancer disorders include, but are not limited to, a solid tumor, a soft tissue tumor (e.g., a lymphoma or a leukemia), and a metastatic lesion. Examples of infectious disorders that can be treated or prevented include bacterial, viral and parasitic infections.

In one embodiment, a method for increasing the ability of a vaccine composition containing an antigen, e.g., an antigen from a pathogen, e.g., a bacterial, viral and parasitic pathogen, or a tumor cell, to elicit a protective immune response in a subject against the antigen by administering to the subject, either simultaneously with or sequentially, to the vaccine composition, an effective adjuvanting amount of an IL-21/IL-21R agonist (e.g., a IL-21 polypeptide or a biologically active fragment thereof, or a nucleic acid encoding the same). In one embodiment, the pathogen against which the vaccine is directed is an intracellular pathogen, e.g., a virus, bacterium, or protozoan. The pathogen may also be an extracellular parasite, e.g., a helminth or bacterium. The antigen may be a whole cell, a protein, a protein subunit or fragment.

Preferred IL-21/IL-21R agonists bind to IL-21 or IL-21R with high affinity, e.g., with an affinity constant of at least about 10^7 M^{-1} , preferably about 10^8 M^{-1} , and more preferably, about 10^9 M^{-1} to 10^{10} M^{-1} or stronger.

In one embodiment, the IL-21/IL-21R agonist is an IL-21 polypeptide, e.g., a human IL-21 polypeptide, or an active fragment thereof (e.g., a human IL-21 polypeptide comprising the amino acid sequence shown as SEQ ID NO:19, or encoded by a nucleotide sequence shown as SEQ ID NO:18, or a sequence at least 85%, 90%, 95%, 98% or more identical thereto). In other embodiments, the IL-21/IL-21R agonist is a fusion protein comprising an IL-21 polypeptide, e.g., human IL-21 polypeptide, or a fragment thereof and, e.g., fused to, a second moiety, e.g., a polypeptide (e.g., an immunoglobulin chain, an Fc fragment, a heavy chain constant regions of the various isotypes, including: IgG1, IgG2, IgG3, IgG4, IgM, IgA1, IgA2, IgD, and IgE); an agonist antibody or antigen-binding fragment thereof, to the IL-21 receptor; or a small molecule or peptide agonist. In other embodiments, the IL-21/IL-21R agonist is an agent that increases the activity or levels of IL-21 by, e.g., increasing expression, processing and/or secretion of functional IL-21. Nucleic acids encoding the aforesaid IL-21/IL-21R agonists and/or antigen can also be administered to the subject.

The IL-21/IL-21R agonists described herein can be used, alone or in combination, with other therapeutic modalities. If desired, the IL-21/IL-21R agonist can be administered in conjunction with one or more additional agents that increase an immune response, e.g., an agent that enhances an immune response to a cancer or infectious disease, in the subject (e.g., an antigen, an antigenic peptide, alone or in combination with an antigen-presenting cell, e.g., a dendritic cell). For example, the IL-21/IL-21R agonist(s) can be administered by themselves, or in combination with an antigen, e.g., as an adjuvant (e.g., a vaccine adjuvant) and/or in combination with other cytokines (e.g., IL-2, GM-CSF and/or IL-15). The combination therapy can be carried out in any order, e.g., IL-15 and antigen can be co-administered to a subject, followed by boosting the immune response by administering IL-21 and the antigen.

Preferably, the subject is a mammal, e.g., a human suffering a cancer or an infectious disorder. The IL-21/IL-21R agonist is preferably designed to increase the immune response to a cancer or infectious disease. The infectious disease can be caused by, e.g., bacterial, parasitic or viral agents.

In another aspect, the invention features a method for modulating, e.g., increasing or decreasing, immune cell activity and/or number (e.g., the activity and/or number of one or more of: a mature T cell (mature, CD8+, CD4+ T cell), mature NK cell, B cell,

macrophage or megakaryocyte; preferably, a mature CD8+ T cell or a macrophage) or a population of immune cells, e.g., a mixed or a substantially purified immune cell population. The method includes contacting an immune cell, e.g., an immune cell as described herein, with an IL-21/IL-21R agonist or antagonist, e.g., an agonist or
5 antagonist as described herein, in an amount sufficient to modulate, e.g., increase or decrease, immune cell activity and/or number.

The subject method can be used on cells in culture, e.g. *in vitro* or *ex vivo*. For example, immune cells, e.g., T cells as described herein, can be cultured *in vitro* in culture medium and the contacting step can be effected by adding one or more IL-21/IL-
10 21R agonist(s) or antagonist(s), e.g., an agonist or antagonist as described herein, to the culture medium. Alternatively, the method can be performed on cells (e.g., immune cells as described herein) present in a subject, e.g., as part of an *in vivo* (e.g., therapeutic or prophylactic) protocol.

The immune cell can be chosen from, e.g., one or more of: a mature T cell
15 (mature, CD8+, CD4+, lymph node T cell, memory T cell), mature NK cell, B cell, antigen presenting cell (APC), e.g., a dendritic cell, macrophage or megakaryocyte, or a population of cells, e.g., a mixed or a substantially purified immune cell population. Preferably, the immune cell is a mature CD8+ T cell or a macrophage.

A change in immune cell activity includes any variation(s), e.g.,
20 increase/decrease, in one or more of: proliferation, cytokine secretion and/or production, survival, differentiation, cell responsiveness (e.g., desensitization), cytolytic activity, effector cell activity, gene expression, among others, of the immune cell contacted with an IL-21/IL-21R agonist or antagonist compared to a reference, e.g., an untreated immune cell. For example, contacting an immune cell with an IL-21/IL-21R agonist,
25 e.g., an IL-21 polypeptide, can induce one or more of: proliferation, cytolytic activity, effector cell function, or cytokine secretion of one or more of: antigen- or anti-CD3 antibody stimulated thymocytes; lymph node T cells, mature CD4+ T cells, mature CD8+ T cells, or macrophages.

Responsiveness of immune cells, e.g., T cells, to stimulatory signals can also be
30 modulated using an agonist or antagonist as described herein. For example, proliferation of T cells to alloantigens can be increased in the presence of an IL-21 polypeptide. IL-21 may also enhance proliferation and/or differentiation of mature CD8+ T cells. For

example, priming of CD8+ T cells in the presence of IL-21 can generate effector cells with enhanced lytic (CTL) activity and/or increased ability to secrete cytokines, e.g., IFN γ . In other embodiments, IL-21/IL-21R agonist can be used to induce the proliferation and/or cytokine secretion of macrophages.

5 In one embodiment, a method for decreasing immune cell activity (e.g., the activity of one or more of: a mature T cell (mature, CD8+, CD4+, lymph node T cell, memory T cell), mature NK cell, B cell, antigen presenting cell (APC), e.g., a dendritic cell, macrophage or megakaryocyte, or a population of cells, e.g., a mixed or a substantially purified immune cell population, is provided. The method includes
10 contacting the immune cell with an IL-21/IL-21R antagonist, e.g., an antagonist as described herein, in an amount sufficient to decrease immune cell activity.

 In other embodiments, a method for increasing immune cell activity and/number (e.g., the activity and/or number of one or more of: a mature T cell (mature, CD8+, CD4+, lymph node T cell, memory T cell), mature NK cell, B cell, antigen presenting
15 cell (APC), e.g., a dendritic cell, macrophage or megakaryocyte, or a population of cells, e.g., a mixed or a substantially purified immune cell population, is provided. The method includes contacting an immune cell with an IL-21/IL-21R agonist, e.g., an agonist as described herein, in an amount sufficient to increase immune cell activity.

 In another aspect, the invention features a fusion protein that includes at least a
20 fragment of an IL-21R polypeptide, which is capable of binding an IL-21 polypeptide, e.g., a soluble fragment of an IL-21R (e.g., a fragment of an IL-21R comprising the extracellular domain of murine or human IL-21R; e.g., from about amino acids 1-235, 1-236, 20-235, 20-236 of SEQ ID NO:2 (human), or from about amino acids 1-236, 20-236 of SEQ ID NO:10 (murine), or encoded by the corresponding nucleotides of SEQ ID
25 NO:1 or 9, or a sequence at least 85%, 90%, 95%, 98% or more identical thereto) and, e.g., fused to, a second moiety, e.g., a polypeptide (e.g., an immunoglobulin chain, an Fc fragment, a heavy chain constant regions of the various isotypes, including: IgG1, IgG2, IgG3, IgG4, IgM, IgA1, IgA2, IgD, and IgE). For example, the fusion protein can include the extracellular domain of human IL-21R, e.g., from about amino acids 1-235,
30 1-236, 20-235, 20-236 of SEQ ID NO:2, and, e.g., fused to, a human immunoglobulin Fc chain (e.g., human IgG, e.g., human IgG1 or a mutated form of human IgG1). In one embodiment, the human Fc sequence has been mutated at one or more amino acids, e.g.,

mutated at residues 254 and 257 of SEQ ID NO:28, from the wild type sequence to reduce Fc receptor binding. In other embodiments, the fusion protein can include the extracellular domain of murine IL-21R, e.g., from about amino acids 1-236, 20-236 of SEQ ID NO:10 (murine), and, e.g., fused to, a murine immunoglobulin Fc chain (e.g.,
5 murine IgG, e.g., murine IgG2a or a mutated form of murine IgG2a). The fusion proteins may additionally include a linker sequence joining the IL-21R fragment to the second moiety. In other embodiments, additional amino acid sequences can be added to the N- or C-terminus of the fusion protein to facilitate expression, detection and/or isolation or purification.

10 The fusion protein described herein, can be derivatized or linked to another functional molecule, e.g., another peptide or protein (e.g., an Fab' fragment). For example, the fusion protein can be functionally linked (e.g., by chemical coupling, genetic fusion, non-covalent association or otherwise) to one or more other molecular entities, such as an antibody (e.g., a bispecific or a multispecific antibody), toxins,
15 radioisotopes, cytotoxic or cytostatic agents, among others.

Examples of antagonistic fusion proteins that can be used in the methods of the invention are shown in Figures 7-15. In one embodiment, the fusion protein includes an amino acid sequence chosen from, e.g., SEQ ID NO:23, SEQ ID NO:25, SEQ ID NO:27, SEQ ID NO:29, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:35, SEQ ID NO:37, or
20 SEQ ID NO:39, or a sequence at least 85%, 90%, 95%, 98% or more identical thereto. In other embodiments, the fusion protein includes an amino acid sequence encoded by a nucleotide sequence chosen from, e.g., SEQ ID NO:22, SEQ ID NO:24, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:30, SEQ ID NO:32, SEQ ID NO:34, SEQ ID NO:36, or SEQ ID NO:38, or a sequence at least 85%, 90%, 95%, 98% or more identical thereto.

25 Preferred fusion proteins have the amino acid sequence shown as SEQ ID NO:25 or SEQ ID NO:29 (Figures 8A-8C and 10A-10C, respectively), or a sequence at least 85%, 90%, 95%, 98% or more identical thereto. In other embodiments, the fusion protein includes an amino acid sequence encoded by a nucleotide sequence chosen from, e.g., SEQ ID NO:24 or SEQ ID NO:28 (Figures 8A-8C and 10A-10C, respectively), or a sequence at
30 least 85%, 90%, 95%, 98% or more identical thereto. Most preferably, the fusion protein has the amino acid sequence shown as SEQ ID NO:29, or has an amino acid sequence encoded by a nucleotide sequence shown as SEQ ID NO:28 (Figure 10A-10C).

The invention also features nucleic acid sequences that encode the fusion proteins described herein.

In another aspect, the invention features host cells and vectors containing the nucleic acids of the invention. Preferably, the host cell is a eukaryotic cell, e.g., a mammalian cell, an insect cell, a yeast cell, or a prokaryotic cell, e.g., *E. coli*. For example, the mammalian cell can be a cultured cell or a cell line. Exemplary mammalian cells include lymphocytic cell lines (e.g., NSO), Chinese hamster ovary cells (CHO), COS cells, oocyte cells, and cells from a transgenic animal, e.g., mammary epithelial cell. For example, nucleic acids encoding the fusion proteins described herein can be expressed in a transgenic animal. In one embodiment, the nucleic acids are placed under the control of a tissue-specific promoter (e.g., a mammary specific promoter) and the antibody is produced in the transgenic animal. For example, the fusion protein is secreted into the milk of the transgenic animal, such as a transgenic cow, pig, horse, sheep, goat or rodent.

In another aspect, the invention provides a process for producing a fusion protein, e.g., a fusion protein as described herein. The process comprises: (a) growing a culture of the host cell of the present invention in a suitable culture medium; and (b) purifying the fusion protein from the culture. Proteins produced according to these methods are also provided.

In another aspect, the invention provides, compositions, e.g., pharmaceutical compositions, which include a pharmaceutically acceptable carrier and at least one of IL-21/IL-21R agonist or an antagonist as described herein (e.g., a fusion protein described herein). In one embodiment, the compositions, e.g., pharmaceutical compositions, comprise a combination of two or more one of the aforesaid IL-21/IL-21R agonists or antagonists. Combinations of the IL-21/IL-21R agonists or antagonists and a drug, e.g., a therapeutic agent (e.g., one or more cytokine and growth factor inhibitors, immunosuppressants, anti-inflammatory agents, metabolic inhibitors, enzyme inhibitors, and/or cytotoxic or cytostatic agents, as described in more herein) or an antigen, e.g., an antigenic peptide and/or an antigen-presenting cell, are also within the scope of the invention.

In one embodiment, the pharmaceutical composition includes an IL-21/IL-21R antagonist agonist and at least one additional therapeutic agent, in a pharmaceutically-acceptable carrier. Examples of preferred additional therapeutic agents that can be co-formulated in a composition, e.g., a pharmaceutical composition, with one or more IL-21/IL-21R antagonists, include, but are not limited to, one or more of: TNF antagonists (e.g., chimeric, humanized, human or *in vitro* generated antibodies, or antigen-binding fragments thereof, that bind to TNF; soluble fragments of a TNF receptor, e.g., p55 or p75 human TNF receptor or derivatives thereof, e.g., 75 kDTNFR-IgG (75 kD TNF receptor-IgG fusion protein, EnbrelTM), p55 kD TNF receptor-IgG fusion protein; TNF enzyme antagonists, e.g., TNF α converting enzyme (TACE) inhibitors); antagonists of IL-12, IL-15, IL-17, IL-18, IL-22; T cell and B cell depleting agents (e.g., anti-CD4 or anti-CD22 antibodies); small molecule inhibitors, e.g., methotrexate and leflunomide; sirolimus (rapamycin) and analogs thereof, e.g., CCI-779; Cox-2 and cPLA2 inhibitors; NSAIDs; p38 inhibitors, TPL-2, M κ -2 and NF κ b inhibitors; RAGE or soluble RAGE; P-selectin or PSGL-1 inhibitors (e.g., small molecule inhibitors, antibodies thereto, e.g., antibodies to P-selectin); estrogen receptor beta (ERB) agonists or ERB-NF κ b antagonists. Most preferred additional therapeutic agents that can be co-administered and/or co-formulated with one or more IL-21/IL-21R antagonists include one or more of: a soluble fragment of a TNF receptor, e.g., p55 or p75 human TNF receptor or derivatives thereof, e.g., 75 kDTNFR-IgG (75 kD TNF receptor-IgG fusion protein, EnbrelTM), methotrexate, leflunomide, or a sirolimus (rapamycin) or an analog thereof, e.g., CCI-779.

In another embodiment, a pharmaceutical composition useful as a vaccine comprising an antigen from a pathogenic microorganism, e.g., a viral, bacterial or parasitic microorganism, and an effective adjuvanting amount of an IL-21/IL-21R agonist, in a pharmaceutically acceptable carrier, is provided. In one embodiment, the resulting composition is capable of eliciting the vaccinated subject's immunity for a protective response to the pathogen. The IL-21/IL-21R agonist and antigen used in the composition may be a polypeptide or biologically active fragment thereof, or in a composition comprising nucleic acids encoding the same. These nucleic acids, together with the appropriate promoter sequences, may be employed directly as an antigen administered with, or close in time to, the IL-21/IL-21R agonist adjuvant. Alternatively,

these nucleic acids sequences may be transduced in alternate vaccine strains of the pathogenic microorganism, and upon expression *in vivo* may provide the antigen of the vaccine.

In other embodiments, the invention provides a composition, e.g., a pharmaceutical composition, for the treatment or amelioration of a cancer, and a method for adjuvanting a cancer "vaccine," in a pharmaceutically acceptable carrier. A cancer vaccine may comprise an antigen expressed on the surface of a cancer or a tumor cell. This antigen may be naturally present on the cancer cell. Alternatively, the cancer cell may be manipulated *ex vivo* and transfected with a selected antigen, which it then expresses when introduced into the patient. An exemplary pharmaceutical composition described herein can contain a cancer- or a tumor cell-antigen (either alone as a protein, biologically active fragment thereof, or nucleic acid encoding same), or a cell, e.g., a cancer cell, transfected with the cancer or tumor cell antigen, in combination with an IL-21/IL-21R agonist (e.g., an agonist as described herein, a fragment thereof, or a nucleic acid encoding the same). In one embodiment, the co-administration of IL-21/IL-21R agonist with the tumor cell antigen enhances the CD8+ T-cell effector cell activity of the tumor cell antigen.

Methods of producing the aforesaid compositions, e.g., vaccine compositions, are also encompassed by the present invention.

The following terms are used interchangeably herein: "MU-1" and "IL-21R," and peptides, polypeptides and proteins.

Unless otherwise defined, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this invention belongs. Although methods and materials similar or equivalent to those described herein can be used in the practice or testing of the invention, suitable methods and materials are described below. All publications, patent applications, patents, and other references mentioned herein are incorporated by reference in their entirety. In the case of conflict, the present specification, including definitions, will control. In addition, the materials, methods, and examples are illustrative only and not intended to be limiting.

Other features and advantages of the invention will be apparent from the following detailed description and claims.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1A depicts the full-length cDNA sequence of murine IL-21R/MU-1. The nucleotide sequence corresponds to nucleotides 1-2628 of SEQ ID NO:9.

Figures 2A-2B depict the amino acid sequences of murine and human IL-21R/MU-1. *Figure 2A* depicts the amino acid sequence of murine IL-21R/MU-1 (corresponding to the amino acids 1-529 of SEQ ID NO:10). There is a predicted leader sequence at amino acids 1-19, which was predicted by Susan with score of 10.1 (bold-face type). There is a predicted transmembrane domain at amino acids 237-253 of SEQ ID NO:10 (underlined). Predicted signaling motifs include the following regions: Box 1: amino acids 265-274 and Box 2: amino acids 310-324 (bold and underlined); six tyrosine's are located at amino acid positions 281, 319, 361, 368, 397, and 510, of SEQ ID NO:10. The WSXWS motif (SEQ ID NO:8) is located at amino acid residue 214 to amino acid residue 218 (in large, bold-face type). Potential STAT docking sites include, amino acids 393-398 and amino acids 510-513 of SEQ ID NO:10.

Figure 2B depicts the amino acid sequence of human MU-1 (corresponding to SEQ ID NO:2). The location of the predicted signal sequence (about amino acids 1-19 of SEQ ID NO:2); WSXWS motif (about amino acids 213-217 of SEQ ID NO:2); and transmembrane domain (about amino acids 236-252, 236-253, 236-254, of SEQ ID NO:2 (underlined)). Potential JAK binding sites, signaling motifs and STAT docking sites are also indicated. The approximate location of these sites is boxed.

Figure 3 depicts the GAP comparison of human and murine MU-1cDNA sequences (corresponding to nucleic acids 1-2665 of SEQ ID NO:1 and nucleic acids 1-2628 of SEQ ID NO:9, respectively). HuMU-1= human MU-1, murMU-1= murine MU-1. Gap Parameters: Gap Weight= 50, Average Match= 10.000, Length Weight= 3, Average Mismatch= 0.000. Percent Identity= 66.116.

Figure 4 depicts a GAP comparison of the human MU-1 protein (corresponding to amino acids 538 of SEQ ID NO:2) and the murine MU-1 protein (corresponding to amino acids 1-529 of SEQ ID NO:10). BLOSUM62 amino acid substitution matrix. (Henikoff, S. and Henikoff, J. G. (1992)). Amino acid substitution matrices from protein blocks (Proc. Natl. Acad. Sci. USA 89: 10915-10919). Gap parameters= Gap Weight: 8,

Average Match= 2.9 12, Length Weight= 2, Average Mismatch= -2.003. Percent Identity= 65.267.

5 *Figure 5* depicts a multiple sequence alignment of the amino acids of human MU-1 (corresponding to SEQ ID NO:2), murine MU-1 (corresponding to SEQ ID NO: 10), and human IL2beta chain (GENbank Accession No. M26062). Leader and transmembrane domains are underlined. Conserved cytokine receptor module motifs are indicated by bold-face type. Potential signaling regions are indicated by underlining and bold-face type.

10 *Figure 6* depicts signaling through MU-1. MU-1 phosphorylates STAT 5 in Clone E7 EPO-MU-1 chimera. Under the conditions specified in Example 3, signaling through MU-1 results in the phosphorylation of STAT 5 at all time-points tested. Treatment of controls or the chimeric BAF-3 cells with IL-3 resulted in phosphorylation of STAT 3, but not STAT 1 or 5.

15 *Figures 7A-7B* depict an alignment of the nucleotide and amino acid sequences of human IL-21R monomer (corresponding to amino acids 20-235 of SEQ ID NO:2) fused at the amino terminal to honey bee leader sequence and His6 tags (amino acids 1-44 of SEQ ID NO:23). The nucleotide and amino acid sequences are shown as SEQ ID NO:22 and SEQ ID NO:23, respectively.

20 *Figures 8A-8C* depict an alignment of the nucleotide and amino acid sequences of human IL-21R extracellular domain (corresponding to amino acids 1-235 of SEQ ID NO:2) fused at the C-terminus via a linker (corresponding to amino acids 236-243 of SEQ ID NO:25) to human immunoglobulin G1 (IgG1) Fc sequence (corresponding to amino acids 244-467 of SEQ ID NO:25). The nucleotide and amino acid sequences are shown as SEQ ID NO:24 and SEQ ID NO:25, respectively.

25 *Figures 9A-9C* depict an alignment of the nucleotide and amino acid sequences of human IL-21R extracellular domain (corresponding to amino acids 1-235 of SEQ ID NO:2) fused at the C-terminus via a linker (corresponding to amino acids 236-243 of SEQ ID NO:27) to human immunoglobulin G1 (IgG1) Fc sequence (corresponding to amino acids 244-467 of SEQ ID NO:27), and His6 sequence tag (corresponding to amino acids 468-492 of SEQ ID NO:27). The nucleotide and amino acid sequences are shown as SEQ ID NO:26 and SEQ ID NO:27, respectively.

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Figures 10A-10C depicts an alignment of the nucleotide and amino acid sequences of human IL-21R extracellular domain (corresponding to amino acids 1-235 of SEQ ID NO:2) fused at the C-terminus via a linker (corresponding to amino acids 236-243 of SEQ ID NO:29) to human immunoglobulin G1 (IgG1) Fc sequence (corresponding to amino acids 244-467 of SEQ ID NO:29) to human immunoglobulin G1 (IgG1) Fc mutated sequence. The human Fc sequence has been mutated at residues 254 and 257 from the wild type sequence to reduce Fc receptor binding. The nucleotide and amino acid sequences are shown as SEQ ID NO:28 and SEQ ID NO:29, respectively.

Figures 11A-11B depict an alignment of the nucleotide and amino acid sequences of human IL-21R extracellular domain (corresponding to amino acids 1-235 of SEQ ID NO:2) fused at the C-terminus to a rhodopsin sequence tag. The nucleotide and amino acid sequences are shown as SEQ ID NO:30 and SEQ ID NO:31, respectively.

Figures 12A-12C depict an alignment of the nucleotide and amino acid sequences of human IL-21R extracellular domain (corresponding to amino acids 1-235 of SEQ ID NO:2) fused at the C-terminus to an EK cleavage site and mutated IgG1 Fc region (corresponding to amino acids 236-470 of SEQ ID NO:33). The nucleotide and amino acid sequences are shown as SEQ ID NO:32 and SEQ ID NO:33, respectively.

Figures 13A-13B depict an alignment of the nucleotide and amino acid sequences of murine IL-21R extracellular domain fused at the C-terminus to mouse immunoglobulin G2a (IgG2a). The nucleotide (genomic) and amino acid sequences are shown as SEQ ID NO:34 and SEQ ID NO:35, respectively.

Figures 14A-14B depict an alignment of the nucleotide and amino acid sequences of murine IL-21R extracellular domain fused at the C-terminus to Flag and His6 sequence tags. The nucleotide (genomic) and amino acid sequences are shown as SEQ ID NO:36 and SEQ ID NO:37, respectively.

Figures 15A-15B depict an alignment of the nucleotide and amino acid sequences of (honey bee leader) murine IL-21R extracellular domain fused at the C-terminus to Flag and His6 sequence tags. The nucleotide (genomic) and amino acid sequences are shown as SEQ ID NO:38 and SEQ ID NO:39, respectively.

Figure 16 is a timetable summarizing the prophylactic, therapeutic and semi-therapeutic treatment schedules for the experiments using collagen-induced arthritis (CIA) mouse models.

Figure 17 is a graph depicting the effects of muIL21RFc (200 µg/mouse 3x/week) on a semi-therapeutic CIA mouse as a function of days post treatment. Mouse Ig (200 µg/mouse 3x/week) was used as a control.

Figure 18 is a graph depicting the effects of murine IL-21 on mean CIA score as a function of days post collagen boost. Mice were injected with 1 µg of murine IL21 intraperitoneally when presenting a total body score of 2-4.

Figure 19, panels A-B, are photographs showing increased expression of IL-21R mRNA in arthritic paws of mice with CIA (panel A) compared to negative controls (panel B).

Figures 20A-20C are bar graphs depicting the level and biological activity of IL-21 secreted by IL-21- transduced B16F1 and MethA cells. Figure 20A shows the level of IL-21 secretion by transduced tumor cells. Figures 20(B, C) are bar graphs depicting the biological activity of IL-21 secreted by transduced tumor cells. Naïve splenocytes from either C57BL/6 (B) or Balb/C (C) mice were stimulated for 72 hours with the indicated concentrations of irradiated syngeneic tumor cells. The cultures were supplemented with sub-optimal amount of anti-CD3 and anti-CD28 mAb in 96-well plates. ³H thymidine was added during the last 6 hours of culture.

Figures 21A-21C are graphs depicting the *in vitro* growth characteristics of transduced tumor cells and evaluation of IL-21R expression. Figures 21A and 21B depict the number of B16F1-IL-21 and B16F1-GFP or MethA-IL-21 and MethA-GFP tumor cells, respectively, with respect to time in culture. Figure 21C is a bar graph depicting expression of IL-21RmRNA in the transfected cells was normalized to cyclophilin values and expressed as relative unite (R.U.).

Figures 22A-22B are linear graphs depicting the *in vivo* growth characteristics of transduced tumor cells. IL-21 or GFP expressing tumor cells were injected in syngeneic naïve mice as indicated. Tumor size was scored by multiplying perpendicular diameters and was averaged for all mice in each group. (A) Tumor growth in C57BL/6 mice that were injected with 10⁵ of B16F1-IL-21 or GFP cells. (B) Tumor growth in Balb/C mice that were injected with either 1x10⁶ or 2x10⁶ of MethA-IL-21 or MethA-GFP cells.

Figures 23A-23B are linear graphs depicting the changes in B16F1-IL-21 tumor growth in scid and nude mice with respect to number of days after injection of B16F1-IL-

21 cells. B16F1-IL-21 and B16F1-GFP tumor cells (10^5 /mouse) were injected into C57BL/6 scid (A) or C57BL/6 nude (B) mice. Tumor size was monitored twice weekly.

Figures 24A-24B are linear graphs depicting the growth of B16F1-IL-21 cells *in vivo* in CD4⁺, CD8⁺ T or NK cell depleted mice. A) C57BL/6 were depleted of CD4⁺ or CD8⁺ T cells by three consecutive injections of anti-CD4 or anti-CD8 mAb on days -3, -2 and -1 before tumor cell injection and every other day post 10^5 B16F1-IL-21 and B16F1-GFP tumor cell injection. (rat IgG was used as isotype control). B) C57BL/6 were depleted of NK cells by one injection of anti-asialo GM1 Ab on day -1 before tumor cell injection and twice per week post 10^5 B16F1-IL-21 and B16F1-GFP tumor cell injection. (rat IgG or rabbit IgG were used as isotype controls for T and NK cell depletion respectively). Tumor size was monitored twice per week as described in Figure 23.

Figures 25A-25C are graphs showing TRP-2 specific T cell responses in B16F1-IL-21 injected mice. (A) Equal numbers of splenocytes ($2 - 4 \times 10^5$) from mice injected with either B16F1-IL-21 or B16F1-GFP were stimulated with $5 \mu\text{g/ml}$ of TRP-2 peptide in the presence of 20 U of IL-2 in an ELISPOT plate pre-coated with anti-IFN γ Ab. After 24 hours, the plate was developed and spot forming units were counted. Results are expressed as number of spot forming units/million of splenocytes (SPU/million splenocytes). (B) Cytolytic activity of splenocytes from B16F1-IL-21 or (C) control B16F1-GFP injected mice were tested against RMA-S cells pulsed with TRP-2 or OVA peptide (control). Cytolytic activity was measured by standard 4-hr Cr⁵¹ Release Assay

Figures 26A-26B are linear graphs depicting the *in vivo* growth of B16F1-IL-21 cells in IFN γ ^{-/-} and IL-10^{-/-} mice. (A) IFN γ ^{-/-} mice or (B) IL-10^{-/-} mice were injected with 10^5 of either B16F1-IL-21 or B16F1-GFP tumor cells and the tumor growth was monitored twice weekly.

Figures 27A-27B are bar graphs depicting IL-21R and IL-21 expression by naïve CD8⁺ T cells. CD8⁺/CD62L⁺ T cells were sorted from LN of 2C TCR Tg mice and stimulated with anti-CD3 ($2.5 \mu\text{g/ml}$ plate-bound), IL-21 (12.5 ng/ml) or anti-CD28 ($7.5 \mu\text{g/ml}$ plate-bound). At the indicated times, cells were harvested, RNA prepared and RPA analysis performed. Samples were quantitated by phosphorimager analysis relative to an internal control. (A) Quantitation of RPA pixels for IL-21R bands. (B) Quantitation of RPA pixels for IL-21 bands.

Figures 28A-28D are linear graphs depicting the effects of IL-21 on proliferation of CD8+ T cells. (A) IL-21 enhances antigen stimulation of CD8+ T cells. Purified 2C CD8+ T cells were stimulated for 72hrs with irradiated Balb/c splenocytes at 5:1 APC:T ratio with the indicated amount of IL-21 or equivalent dilution of mock-transfected supernatant (representative experiment, n=5). (B) Effect of IL-21 vs IL-2, IL-12, IL-15 on antigen stimulation. 2C T cells were stimulated with 3:1 T-depleted APC:T ratio and the indicated concentration of cytokine (n=3). (C) IL-21 acts directly on CD8+ T cells to specifically enhance proliferation. 2C CD8+ T cells were stimulated on anti-CD3-coated plates (2.5ug/ml) with the indicated titer of IL-21 +/- mIL-21R.Fc (20ug/ml) (n=3). (D) Effect of IL-21 vs IL-2, IL-12, IL-15 on anti-CD3 mAb stimulation (5ug/ml plate-bound).

Figures 29A-29C are graphs showing the effects of IL-21 on development of CD8+ T cell effector functions. (A) IL-21 enhances CTL generation. 2C T cells were stimulated with irradiated APCs (APC:T) and IL-2 (10U/ml), IL-21 (titer) or the equivalent dilution of mock supernatant for 4 days then washed, counted and set up in a 4hr CTL assays using 51-Cr-labeled P815 as specific targets (representative experiment, n=3). (B) IL-21 is comparable to IL-12, IL-15 for inducing lytic activity from CD8+ T cells. 2C CD8+ T cells were stimulated and assayed for CTL activity as in (A) using IL-2 (5ng/ml), IL-15 (50ng/ml), IL-12 (5ng/ml), IL-21 (25ng/ml). (C) IL-21 during priming results in increased ability to produce IFNg. 2C T cells were stimulated as indicated in (B) for 5 days then washed and restimulated for on anti-CD3-coated plates (1ug/ml). IFNg ELISAs were done on 40hr supernatants.

DETAILED DESCRIPTION OF THE INVENTION

Methods and compositions for modulating interleukin-21 (IL-21)/IL-21 receptor (MU-1) activity using agonists or antagonists of IL-21 or IL-21 receptor ("IL-21R" or "MU-1"), are disclosed. IL-21/IL-21R antagonists can be used to induce immune suppression *in vivo*, e.g., for treating or preventing immune cell-associated pathologies (e.g., pathologies associated with aberrant activity of one or more of mature T cells (mature CD8+, mature CD4+ T cells), mature NK cells, B cells, macrophages and megakaryocytes, including transplant rejection and autoimmune disorders). IL-21/IL-21R agonists can be used by themselves or in combination with an antigen, e.g., as an adjuvant (e.g., a vaccine adjuvant), to up-regulate an immune response *in vivo*, e.g., for example, for use in treating cancer and infectious disorders.

In one embodiment, Applicants have shown that a reduction of IL-21R activity by using a neutralizing fusion protein that includes the extracellular domain of the IL-21R fused to an Fc immunoglobulin region ameliorates inflammatory symptoms in mouse collagen-induced arthritis (CIA) animal models (Example 7). Expression of IL-21R mRNA is upregulated in the paws of CIA mice (Example 8). Accordingly, IL-21R binding agents that antagonize IL-21/IL-21R activity can be used to induce immune suppression *in vivo*, e.g., for treating or preventing immune cell-associated pathologies (e.g., pathologies associated with aberrant activity of one or more of T cells (CD8+, CD4+ T cells), NK cells, B cells, macrophages and megakaryocytes, including transplant rejection and autoimmune disorders).

In other embodiments, Applicants have shown that agonistic IL-21R binding agent stimulate, primarily via stimulation of CD8+ T cells, innate and adaptive immunity *in vivo* against immunogenic and non-immunogenic tumor cells (Example 9). Accordingly, binding agents that stimulate the IL-21/IL-21R pathway can be used by themselves or in combination with an antigen, e.g., as an adjuvant (e.g., a vaccine adjuvant), to up-regulate an immune response *in vivo*, e.g., for example, for use in treating cancer and infectious disorders.

In order that the present invention may be more readily understood, certain terms are first defined. Additional definitions are set forth throughout the detailed description.

The term "MU-1," "MU-1 protein," "interleukin-21 receptor" or "IL-21R," as used herein, refers to a class I cytokine family receptor, also known as NILR (WO 01/85792; Parrish-Novak *et al.* (2000) *Nature* 408:57-63; Ozaki *et al.* (2000) *Proc. Natl. Acad. Sci. USA* 97:11439-114444). MU-1 is homologous to the shared β chain of the IL-2 and IL-15 receptors, and IL-4 α (Ozaki *et al.* (2000) *supra*). Upon ligand binding, IL-21R/MU-1 is capable of interacting with a common γ cytokine receptor chain (γ_c) (Asao *et al.* (2001) *J. Immunol.* 167:1-5), and inducing the phosphorylation of STAT1 and STAT3 (Asao *et al.* (2001) or STAT5 (Ozaki *et al.* (2000). MU-1 shows widespread lymphoid tissue distribution. The term "MU-1" refers to a receptor (preferably of mammalian, e.g., murine or human origin) which is capable of interacting with, e.g., binding to, IL-21 (preferably of mammalian, e.g., murine or human IL-21) and having one of the following features: (i) an amino acid sequence of a naturally occurring mammalian MU-1 polypeptide IL-21R/MU-1 or a fragment thereof, e.g., an amino acid

sequence shown as SEQ ID NO:2 (human) or SEQ ID NO:10 (murine) or a fragment thereof; (ii) an amino acid sequence substantially homologous to, e.g., at least 85%, 90%, 95%, 98%, 99% homologous to, an amino acid sequence shown as SEQ ID NO:2 (human) or SEQ ID NO:10 (murine) or a fragment thereof; (iii) an amino acid sequence
5 which is encoded by a naturally occurring mammalian IL-21R/MU-1 nucleotide sequence or a fragment thereof (e.g., SEQ ID NO:1 (human) or SEQ ID NO:9 (murine) or a fragment thereof); (iv) an amino acid sequence encoded by a nucleotide sequence which is substantially homologous to, e.g., at least 85%, 90%, 95%, 98%, 99% homologous to, a nucleotide sequence shown as SEQ ID NO:1 (human) or SEQ ID NO:9 (murine) or a
10 fragment thereof; (v) an amino acid sequence encoded by a nucleotide sequence degenerate to a naturally occurring IL-21R/MU-1 nucleotide sequence or a fragment thereof, e.g., SEQ ID NO:1 (human) or SEQ ID NO:9 (murine) or a fragment thereof; or (vi) a nucleotide sequence that hybridizes to one of the foregoing nucleotide sequence sequences under stringent conditions, e.g., highly stringent conditions.

15 The IL-21R/MU-1 is of mammalian, preferably, human origin. The nucleotide sequence and the predicted amino acid sequence of human IL-21R/MU-1 are shown in SEQ ID NO:1 and SEQ ID NO:2, respectively. Analysis of the human IL-21R/MU-1 amino acid sequence (SEQ ID NO:2; Figure 2B) revealed the following structural features: a leader sequence from about (about amino acids 1-19 of SEQ ID NO:2 (Figure
20 2B)); WSXWS motif (about amino acids 213-217 of SEQ ID NO:2); transmembrane domain (about amino acids 236-252 of SEQ ID NO:2 (Figure 2B)); an extracellular domain from about amino acids 1-235 of SEQ ID NO:2; and an intracellular domain from about 253-538 of SEQ ID NO:2. The mature human IL-21R/MU-1 is believed to have the sequence of amino acids 20-538 of SEQ ID NO:2.

25 The IL-21R/MU-1 cDNA was deposited with the American Type Culture Collection on Mar. 10, 1998, as accession number ATCC 98687.

Any form of IL-21R/MU-1 proteins of less than full length can be used in the methods and compositions of the present invention, provided that it retains the ability to bind to an IL-21 polypeptide. IL-21R/MU-1 proteins of less than full length, e.g., soluble
30 IL-21R, can be produced by expressing a corresponding fragment of the polynucleotide encoding the full-length MU-1 protein in a host-cell. These corresponding polynucleotide fragments are also part of the present invention. Modified

polynucleotides as described above may be made by standard molecular biology techniques, including construction of appropriate desired deletion mutants, site-directed mutagenesis methods or by the polymerase chain reaction using appropriate oligonucleotide primers.

5 As used herein, a "soluble IL-21R/MU-1 polypeptide" is a IL-21R/MU-1 polypeptide incapable of anchoring itself in a membrane. Such soluble polypeptides include, for example, MU-1 or IL-21R polypeptides that lack a sufficient portion of their membrane spanning domain to anchor the polypeptide or are modified such that the membrane spanning domain is non-functional. E.g., a soluble fragment of an IL-21R
10 (e.g., a fragment of an IL-21R comprising the extracellular domain of murine or human IL-21R includes an amino acid sequence from about amino acids 1-235, 1-236, 20-235, 20-236 of SEQ ID NO:2 (human), or from about amino acids 1-236, 20-236 of SEQ ID NO:10 (murine). A soluble IL-21R/MU-1 polypeptide can additionally include, e.g., be fused to, a second moiety, e.g., a polypeptide (e.g., an immunoglobulin chain, a GST,
15 Lex-A or MBP polypeptide sequence). For example, a fusion protein can includes at least a fragment of an IL-21R polypeptide, which is capable of binding IL-21, e.g., a soluble fragment of an IL-21R (e.g., a fragment of an IL-21R comprising the extracellular domain of murine or human IL-21R; e.g., from about amino acids acids 1-235, 1-236, 20-235, 20-236 of SEQ ID NO:2 (human), or from about amino acids 1-236, 20-236 of SEQ
20 ID NO:10 (murine), fused to a second moiety, e.g., a polypeptide (e.g., an immunoglobulin chain, an Fc fragment, a heavy chain constant regions of the various isotypes, including: IgG1, IgG2, IgG3, IgG4, IgM, IgA1, IgA2, IgD, and IgE).

The term "interleukin-21" or "IL-21" refers to a cytokine showing sequence homology to IL-2, IL-4 and IL-15 (Parrish-Novak *et al.* (2000) *Nature* 408:57-63).
25 Despite low sequence homology among interleukin cytokines, cytokines share a common fold into a "four-helix-bundle" structure that is representative of the family. It is expressed primarily in activated CD4+ T cells, and has been reported to have effects on NK, B and T cells (Parrish-Novak *et al.* (2000) *supra*; Kasaian, M.T. *et al.* (2002) *supra*). IL-21 binds to IL-21R (also referred to herein as MU-1 and NILR). Upon IL-21 binding,
30 activation of IL-21R leads to stat5 or stat3 signaling (Ozaki *et al.* (2000) *supra*). The term "IL-21" or "IL-21 polypeptide" refers to a protein (preferably of mammalian, e.g., murine or human origin) which is capable of interacting with, e.g., binding to, IL-21R

(preferably of mammalian, e.g., murine or human IL-21) and having one of the following features: (i) an amino acid sequence of a naturally occurring mammalian IL-21 or a fragment thereof, e.g., an amino acid sequence shown as SEQ ID NO:19 (human) or a fragment thereof; (ii) an amino acid sequence substantially homologous to, e.g., at least 85%, 90%, 95%, 98%, 99% homologous to, an amino acid sequence shown as SEQ ID NO:19 (human) or a fragment thereof; (iii) an amino acid sequence which is encoded by a naturally occurring mammalian IL-21 nucleotide sequence or a fragment thereof (e.g., SEQ ID NO:18 (human) or a fragment thereof); (iv) an amino acid sequence encoded by a nucleotide sequence which is substantially homologous to, e.g., at least 85%, 90%, 95%, 98%, 99% homologous to, a nucleotide sequence shown as SEQ ID NO:18 (human) or a fragment thereof; (v) an amino acid sequence encoded by a nucleotide sequence degenerate to a naturally occurring IL-21 nucleotide sequence or a fragment thereof, e.g., SEQ ID NO:19 (human) or a fragment thereof; or (vi) a nucleotide sequence that hybridizes to one of the foregoing nucleotide sequence sequences under stringent conditions, e.g., highly stringent conditions.

The phrase "a biological activity of" a MU-1 or IL-21R polypeptide refers to one or more of the biological activities of the corresponding mature MU-1 protein, including, but not limited to, (1) interacting with, e.g., binding to, an IL-21 polypeptide; (2) associating with signal transduction molecules, e.g., γ c, jak1; (3) stimulating phosphorylation and/or activation of stat proteins, e.g., stat 5 and/or stat3; and/or (4) modulating, e.g., stimulating or decreasing, proliferation, differentiation, effector cell function, cytolytic activity, cytokine secretion, and/or survival of immune cells, e.g., T cells (CD8+, CD4+ T cells), NK cells, B cells, macrophages and megakaryocytes).

As used herein, a "IL-21/IL-21R agonist", that is useful in the method of the invention, refers to an agent which potentiates, induces or otherwise enhances one or biological activities of an IL-21R/MU-1 polypeptide. Preferably, the agonist interacts with, e.g., binds to, an IL-21R/MU-1 polypeptide.

As used herein, a "IL-21/IL-21R antagonist", that is useful in the method of the invention, refers to an agent which reduces, inhibits or otherwise diminishes one or biological activities of an IL-21R/MU-1 polypeptide. Preferably, the antagonist interacts with, e.g., binds to, an IL-21R/MU-1 polypeptide. Antagonism using an IL-21/IL-21R

antagonist does not necessarily indicate a total elimination of the IL-21R/MU-1 polypeptide biological activity.

As used herein, a "therapeutically effective amount" of an IL-21/IL-21R agonist or antagonist refers to an amount of an agent which is effective, upon single or multiple
5 dose administration to a subject, e.g., a human patient, at curing, reducing the severity of, ameliorating one or more symptoms of a disorder, or in prolonging the survival of the subject beyond that expected in the absence of such treatment.

As used herein, "a prophylactically effective amount" of an IL-21/IL-21R agonist or antagonist refers to an amount of an IL-21/IL-21R agonist or antagonist which is
10 effective, upon single- or multiple-dose administration to a subject, e.g., a human patient, in preventing or delaying the occurrence of the onset or recurrence of a disorder, e.g., a disorder as described herein.

The term "induce", "inhibit", "potentiate", "elevate", "increase", "decrease" or the like, e.g., which denote quantitative differences between two states, refer to at least
15 statistically significant differences between the two states.

The term "in combination" in this context means that the agents are given substantially contemporaneously, either simultaneously or sequentially. If given sequentially, at the onset of administration of the second compound, the first of the two compounds is preferably still detectable at effective concentrations at the site of
20 treatment.

As used herein, a "fusion protein" refers to a protein containing two or more operably associated, e.g., linked, moieties, e.g., protein moieties. Preferably, the moieties are covalently associated. The moieties can be directly associate, or connected via a spacer or linker.

As used herein, the term "antibody" refers to a protein comprising at least one, and preferably two, heavy (H) chain variable regions (abbreviated herein as VH), and at least one and preferably two light (L) chain variable regions (abbreviated herein as VL). The VH and VL regions can be further subdivided into regions of hypervariability, termed "complementarity determining regions" ("CDR"), interspersed with regions that
30 are more conserved, termed "framework regions" (FR). The extent of the framework region and CDR's has been precisely defined (see, Kabat, E.A., *et al.* (1991) *Sequences of Proteins of Immunological Interest, Fifth Edition*, U.S. Department of Health and Human

Services, NIH Publication No. 91-3242, and Chothia, C. et al. (1987) J. Mol. Biol. 196:901-917, which are incorporated herein by reference). Each VH and VL is composed of three CDR's and four FRs, arranged from amino-terminus to carboxy-terminus in the following order: FR1, CDR1, FR2, CDR2, FR3, CDR3, FR4.

5 The antibody can further include a heavy and light chain constant region, to thereby form a heavy and light immunoglobulin chain, respectively. In one embodiment, the antibody is a tetramer of two heavy immunoglobulin chains and two light immunoglobulin chains, wherein the heavy and light immunoglobulin chains are interconnected by, e.g., disulfide bonds. The heavy chain constant region is comprised of
10 three domains, CH1, CH2 and CH3. The light chain constant region is comprised of one domain, CL. The variable region of the heavy and light chains contains a binding domain that interacts with an antigen. The constant regions of the antibodies typically mediate the binding of the antibody to host tissues or factors, including various cells of the immune system (e.g., effector cells) and the first component (C1q) of the classical
15 complement system.

As used herein, the term "immunoglobulin" refers to a protein consisting of one or more polypeptides substantially encoded by immunoglobulin genes. The recognized human immunoglobulin genes include the kappa, lambda, alpha (IgA1 and IgA2), gamma (IgG1, IgG2, IgG3, IgG4), delta, epsilon and mu constant region genes, as well as
20 the myriad immunoglobulin variable region genes. Full-length immunoglobulin "light chains" (about 25 Kd or 214 amino acids) are encoded by a variable region gene at the NH2-terminus (about 110 amino acids) and a kappa or lambda constant region gene at the COOH-terminus. Full-length immunoglobulin "heavy chains" (about 50 Kd or 446 amino acids), are similarly encoded by a variable region gene (about 116 amino acids)
25 and one of the other aforementioned constant region genes, e.g., gamma (encoding about 330 amino acids).

As used herein, "isotype" refers to the antibody class (e.g., IgM or IgG1) that is encoded by heavy chain constant region genes.

The term "antigen-binding fragment" of an antibody (or simply "antibody
30 portion," or "fragment"), as used herein, refers to one or more fragments of a full-length antibody that retain the ability to specifically bind to an antigen (e.g., CD3). Examples of binding fragments encompassed within the term "antigen-binding fragment" of an

antibody include (i) a Fab fragment, a monovalent fragment consisting of the VL, VH, CL and CH1 domains; (ii) a F(ab')₂ fragment, a bivalent fragment comprising two Fab fragments linked by a disulfide bridge at the hinge region; (iii) a Fd fragment consisting of the VH and CH1 domains; (iv) a Fv fragment consisting of the VL and VH domains of a single arm of an antibody, (v) a dAb fragment (Ward *et al.*, (1989) *Nature* 341:544-546), which consists of a VH domain; and (vi) an isolated complementarity determining region (CDR). Furthermore, although the two domains of the Fv fragment, VL and VH, are coded for by separate genes, they can be joined, using recombinant methods, by a synthetic linker that enables them to be made as a single protein chain in which the VL and VH regions pair to form monovalent molecules (known as single chain Fv (scFv); see *e.g.*, Bird *et al.* (1988) *Science* 242:423-426; and Huston *et al.* (1988) *Proc. Natl. Acad. Sci. USA* 85:5879-5883). Such single chain antibodies are also intended to be encompassed within the term "antigen-binding fragment" of an antibody. These antibody fragments are obtained using conventional techniques known to those with skill in the art, and the fragments are screened for utility in the same manner as are intact antibodies.

Sequences similar or homologous (*e.g.*, at least about 85% sequence identity) to the sequences disclosed herein are also part of this application. In some embodiment, the sequence identity can be about 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or higher. Alternatively, substantial identity exists when the nucleic acid segments will hybridize under selective hybridization conditions (*e.g.*, highly stringent hybridization conditions), to the complement of the strand. The nucleic acids may be present in whole cells, in a cell lysate, or in a partially purified or substantially pure form.

Calculations of "homology" or "sequence identity" between two sequences (the terms are used interchangeably herein) are performed as follows. The sequences are aligned for optimal comparison purposes (*e.g.*, gaps can be introduced in one or both of a first and a second amino acid or nucleic acid sequence for optimal alignment and non-homologous sequences can be disregarded for comparison purposes). In a preferred embodiment, the length of a reference sequence aligned for comparison purposes is at least 30%, preferably at least 40%, more preferably at least 50%, even more preferably at least 60%, and even more preferably at least 70%, 80%, 90%, 100% of the length of the reference sequence. The amino acid residues or nucleotides at corresponding amino acid positions or nucleotide positions are then compared. When a position in the first

sequence is occupied by the same amino acid residue or nucleotide as the corresponding position in the second sequence, then the molecules are identical at that position (as used herein amino acid or nucleic acid "identity" is equivalent to amino acid or nucleic acid "homology"). The percent identity between the two sequences is a function of the
5 number of identical positions shared by the sequences, taking into account the number of gaps, and the length of each gap, which need to be introduced for optimal alignment of the two sequences.

The comparison of sequences and determination of percent identity between two sequences can be accomplished using a mathematical algorithm. In a preferred
10 embodiment, the percent identity between two amino acid sequences is determined using the Needleman and Wunsch ((1970) *J. Mol. Biol.* 48:444-453) algorithm which has been incorporated into the GAP program in the GCG software package (available at <http://www.gcg.com>), using either a Blossum 62 matrix or a PAM250 matrix, and a gap weight of 16, 14, 12, 10, 8, 6, or 4 and a length weight of 1, 2, 3, 4, 5, or 6. In yet
15 another preferred embodiment, the percent identity between two nucleotide sequences is determined using the GAP program in the GCG software package (available at <http://www.gcg.com>), using a NWSgapdna.CMP matrix and a gap weight of 40, 50, 60, 70, or 80 and a length weight of 1, 2, 3, 4, 5, or 6. A particularly preferred set of parameters (and the one that should be used if the practitioner is uncertain about what
20 parameters should be applied to determine if a molecule is within a sequence identity or homology limitation of the invention) are a Blossum 62 scoring matrix with a gap penalty of 12, a gap extend penalty of 4, and a frameshift gap penalty of 5. The percent identity between two amino acid or nucleotide sequences can also be determined using
25 the algorithm of E. Meyers and W. Miller ((1989) CABIOS, 4:11-17) which has been incorporated into the ALIGN program (version 2.0), using a PAM120 weight residue table, a gap length penalty of 12 and a gap penalty of 4.

As used herein, the term "hybridizes under stringent conditions" describes conditions for hybridization and washing. Stringent conditions are known to those skilled in the art and can be found in *Current Protocols in Molecular Biology*, John
30 Wiley & Sons, N.Y. (1989), 6.3.1-6.3.6. Aqueous and nonaqueous methods are described in that reference and either can be used. A preferred, example of stringent hybridization conditions are hybridization in 6X sodium chloride/sodium citrate (SSC) at

about 45°C, followed by one or more washes in 0.2X SSC, 0.1% SDS at 50°C. Another example of stringent hybridization conditions are hybridization in 6X SSC at about 45°C, followed by one or more washes in 0.2X SSC, 0.1% SDS at 55°C. A further example of stringent hybridization conditions are hybridization in 6X SSC at about 45°C, followed by one or more washes in 0.2X SSC, 0.1% SDS at 60°C. Preferably, stringent hybridization conditions are hybridization in 6X SSC at about 45°C, followed by one or more washes in 0.2X SSC, 0.1% SDS at 65°C. Particularly preferred highly stringent conditions (and the conditions that should be used if the practitioner is uncertain about what conditions should be applied to determine if a molecule is within a hybridization limitation of the invention) are 0.5M sodium phosphate, 7% SDS at 65°C, followed by one or more washes at 0.2X SSC, 1% SDS at 65°C.

It is understood that the IL-21/IL-21R agonists and antagonists of the present invention may have additional conservative or non-essential amino acid substitutions, which do not have a substantial effect on their functions.

A "conservative amino acid substitution" is one in which the amino acid residue is replaced with an amino acid residue having a similar side chain. Families of amino acid residues having similar side chains have been defined in the art. These families include amino acids with basic side chains (e.g., lysine, arginine, histidine), acidic side chains (e.g., aspartic acid, glutamic acid), uncharged polar side chains (e.g., glycine, asparagine, glutamine, serine, threonine, tyrosine, cysteine), nonpolar side chains (e.g., alanine, valine, leucine, isoleucine, proline, phenylalanine, methionine, tryptophan), beta-branched side chains (e.g., threonine, valine, isoleucine) and aromatic side chains (e.g., tyrosine, phenylalanine, tryptophan, histidine).

The term "recombinant host cell" (or simply "host cell"), as used herein, is intended to refer to a cell into which a recombinant expression vector has been introduced. It should be understood that such terms are intended to refer not only to the particular subject cell, but to the progeny of such a cell. Because certain modifications may occur in succeeding generations due to either mutation or environmental influences, such progeny may not, in fact, be identical to the parent cell, but are still included within the scope of the term "host cell" as used herein.

IL-21/IL-21R Agonists and Antagonists

In one embodiment, an IL-21R/MU-1 polypeptide or active fragments thereof may be fused to a second moiety, e.g., an immunoglobulin or a fragment thereof (e.g., an Fc binding fragment thereof). For example, soluble forms of the IL-21R/MU-1 may be fused through "linker" sequences to the Fc portion of an immunoglobulin. Other fusion proteins, such as those with GST, Lex-A or MBP, may also be used.

The fusion proteins may additionally include a linker sequence joining the IL-21 or IL-21R fragment to the second moiety. For example, the fusion protein can include a peptide linker, e.g., a peptide linker of about 4 to 20, more preferably, 5 to 10, amino acids in length; the peptide linker is 8 amino acids in length. Each of the amino acids in the peptide linker is selected from the group consisting of Gly, Ser, Asn, Thr and Ala; the peptide linker includes a Gly-Ser element. In other embodiments, the fusion protein includes a peptide linker and the peptide linker includes a sequence having the formula (Ser-Gly-Gly-Gly-Gly)_y wherein y is 1, 2, 3, 4, 5, 6, 7, or 8.

In other embodiments, additional amino acid sequences can be added to the N- or C-terminus of the fusion protein to facilitate expression, detection and/or isolation or purification. For example, IL-21/IL-21R fusion protein may be linked to one or more additional moieties, e.g., GST, His6 tag, FLAG tag. For example, the fusion protein may additionally be linked to a GST fusion protein in which the fusion protein sequences are fused to the C-terminus of the GST (*i.e.*, glutathione S-transferase) sequences. Such fusion proteins can facilitate the purification of the MU-1 fusion protein.

In another embodiment, the fusion protein includes a heterologous signal sequence (*i.e.*, a polypeptide sequence that is not present in a polypeptide encoded by a Mu-1 nucleic acid) at its N-terminus. For example, the native Mu-1 signal sequence can be removed and replaced with a signal sequence from another protein. In certain host cells (*e.g.*, mammalian host cells), expression and/or secretion of Mu-1 can be increased through use of a heterologous signal sequence.

A chimeric or fusion protein of the invention can be produced by standard recombinant DNA techniques. For example, DNA fragments coding for the different polypeptide sequences are ligated together in-frame in accordance with conventional techniques, *e.g.*, by employing blunt-ended or stagger-ended termini for ligation, restriction enzyme digestion to provide for appropriate termini, filling-in of cohesive

ends as appropriate, alkaline phosphatase treatment to avoid undesirable joining, and enzymatic ligation. In another embodiment, the fusion gene can be synthesized by conventional techniques including automated DNA synthesizers. Alternatively, PCR amplification of gene fragments can be carried out using anchor primers that give rise to
5 complementary overhangs between two consecutive gene fragments that can subsequently be annealed and reamplified to generate a chimeric gene sequence (see, for example, Ausubel *et al.* (eds.) CURRENT PROTOCOLS IN MOLECULAR BIOLOGY, John Wiley & Sons, 1992). Moreover, many expression vectors are commercially available that encode a fusion moiety (e.g., an Fc region of an immunoglobulin heavy chain). A
10 Mu-1 encoding nucleic acid can be cloned into such an expression vector such that the fusion moiety is linked in-frame to the immunoglobulin protein.

In some embodiments, MU-1 fusion polypeptides exist as oligomers, such as dimers or trimers.

The first polypeptide, and/or nucleic acids encoding the first polypeptide, can be
15 constructed using methods known in the art.

In some embodiments, the Mu-1 polypeptide moiety is provided as a variant Mu-1 polypeptide having a mutation in the naturally-occurring Mu-1 sequence (wild type) that results in higher affinity (relative to the non-mutated sequence) binding of the Mu-1 polypeptide to a IL-21.

20 In some embodiments, the Mu-1 polypeptide moiety is provided as a variant Mu-1 polypeptide having mutations in the naturally-occurring Mu-1 sequence (wild type) that results in a Mu-1 sequence more resistant to proteolysis (relative to the non-mutated sequence).

In some embodiments, the first polypeptide includes full-length Mu-1
25 polypeptide. Alternatively, the first polypeptide comprise less than full-length Mu-1 polypeptide.

A signal peptide that can be included in the fusion protein is
MPLLLLLLLLLPSPLHP (SEQ ID NO:21). If desired, one or more amino acids can
30 additionally be inserted between the first polypeptide moiety comprising the Mu-1 moiety and the second polypeptide moiety.

The second polypeptide is preferably soluble. In some embodiments, the second polypeptide enhances the half-life, (e.g., the serum half-life) of the linked polypeptide. In

some embodiments, the second polypeptide includes a sequence that facilitates association of the fusion polypeptide with a second Mu-1 polypeptide. In preferred embodiments, the second polypeptide includes at least a region of an immunoglobulin polypeptide. Immunoglobulin fusion polypeptides are known in the art and are described
5 in e.g., US Patent Nos. 5,516,964; 5,225,538; 5,428,130; 5,514,582; 5,714,147; and 5,455,165.

In some embodiments, the second polypeptide comprises a full-length immunoglobulin polypeptide. Alternatively, the second polypeptide comprises less than full-length immunoglobulin polypeptide, e.g., a heavy chain, light chain, Fab, Fab₂, Fv,
10 or Fc. Preferably, the second polypeptide includes the heavy chain of an immunoglobulin polypeptide. More preferably, the second polypeptide includes the Fc region of an immunoglobulin polypeptide.

In another aspect of the invention the second polypeptide has less effector function than the effector function of a Fc region of a wild-type immunoglobulin heavy
15 chain. Fc effector function includes for example, Fc receptor binding, complement fixation and T cell depleting activity (*see for example*, US Patent No. 6,136,310). Methods for assaying T cell depleting activity, Fc effector function, and antibody stability are known in the art. In one embodiment the second polypeptide has low or no affinity for the Fc receptor. In an alternative embodiment, the second polypeptide has low or no
20 affinity for complement protein C1q.

A preferred second polypeptide sequence includes the amino acid sequence of SEQ ID NO: 17. This sequence includes a Fc region. Underlined amino acids are those that differ from the amino acid found in the corresponding position of the wild-type immunoglobulin sequence:
25 HTCPPCPAPEALGAPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHN:
K
TKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPVPPIEKTISKAKGQPREPQVYTL
P
30 SREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFFLYSKLTVDKSR'
Q
QGNVFSCSVMHEALHNHYTQKSLSLSPGK (SEQ ID NO:17)

Examples of antagonistic fusion proteins that can be used in the methods of the
35 invention are shown in Figures 7-15. In one embodiment, the fusion protein includes an amino acid sequence chosen from, e.g., SEQ ID NO:23, SEQ ID NO:25, SEQ ID NO:27,

SEQ ID NO:29, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:35, SEQ ID NO:37, or SEQ ID NO:39, or a sequence at least 85%, 90%, 95%, 98% or more identical thereto. In other embodiments, the fusion protein includes an amino acid sequence encoded by a nucleotide sequence chosen from, e.g., SEQ ID NO:22, SEQ ID NO:24, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:30, SEQ ID NO:32, SEQ ID NO:34, SEQ ID NO:36, or SEQ ID NO:38, or a sequence at least 85%, 90%, 95%, 98% or more identical thereto. Preferred fusion proteins have the amino acid sequence shown as SEQ ID NO:25 or SEQ ID NO:29 (Figures 8A-8C and 10A-10C, respectively), or a sequence at least 85%, 90%, 95%, 98% or more identical thereto. In other embodiments, the fusion protein includes an amino acid sequence encoded by a nucleotide sequence chosen from, e.g., SEQ ID NO:24 or SEQ ID NO:28 (Figures 8A-8C and 10A-10C, respectively), or a sequence at least 85%, 90%, 95%, 98% or more identical thereto. Most preferably, the fusion protein has the amino acid sequence shown as SEQ ID NO:29, or has an amino acid sequence encoded by a nucleotide sequence shown as SEQ ID NO:28 (Figure 10A-10C).

In other embodiments, the IL-21/IL-21R agonists or antagonists are antibodies, or antigen-binding fragments thereof, that bind to IL-21 or IL-21R, preferably, mammalian (e.g., human or murine) IL-21 or IL-21R.

MU-1 proteins of the invention may also be used to immunize animals to obtain polyclonal and monoclonal antibodies which specifically react with the MU-1 protein and which may inhibit binding of ligands to the receptor. Such antibodies may be obtained using the entire MU-1 as an immunogen, or by using fragments of MU-1. Smaller fragments of the MU-1 may also be used to immunize animals. The peptide immunogens additionally may contain a cysteine residue at the carboxyl terminus, and are conjugated to a hapten such as keyhole limpet hemocyanin (KLH). Additional peptide immunogens may be generated by replacing tyrosine residues with sulfated tyrosine residues. Methods for synthesizing such peptides are known in the art, for example, as in R. P. Merrifield, J. Amer. Chem. Soc. 85, 2149-2154 (1963); J. L. Krstenansky, et al., FEBS Lett. 211, 10 (1987).

Neutralizing or non-neutralizing antibodies (preferably monoclonal antibodies) binding to MU-1 protein may also be useful in the treatment of conditions described above. These neutralizing monoclonal antibodies may be capable of blocking ligand binding to the MU-1 receptor chain.

The present invention further provides for compositions comprising an antibody that specifically reacts with an IL-21 or an IL-21R.

Human monoclonal antibodies (mAbs) directed against IL-21 or IL-21R can be generated using transgenic mice carrying the human immunoglobulin genes rather than the mouse system. Splenocytes from these transgenic mice immunized with the antigen of interest are used to produce hybridomas that secrete human mAbs with specific affinities for epitopes from a human protein (see, e.g., Wood et al. International Application WO 91/00906, Kucherlapati et al. PCT publication WO 91/10741; Lonberg et al. International Application WO 92/03918; Kay et al. International Application WO 92/03917; Lonberg, N. et al. 1994 *Nature* 368:856-859; Green, L.L. et al. 1994 *Nature Genet.* 7:13-21; Morrison, S.L. et al. 1994 *Proc. Natl. Acad. Sci. USA* 81:6851-6855; Bruggeman et al. 1993 *Year Immunol* 7:33-40; Tuaillon et al. 1993 *PNAS* 90:3720-3724; Bruggeman et al. 1991 *Eur J Immunol* 21:1323-1326).

Monoclonal antibodies can also be generated by other methods known to those skilled in the art of recombinant DNA technology. An alternative method, referred to as the "combinatorial antibody display" method, has been developed to identify and isolate antibody fragments having a particular antigen specificity, and can be utilized to produce monoclonal antibodies (for descriptions of combinatorial antibody display see e.g., Sastry et al. 1989 *PNAS* 86:5728; Huse et al. 1989 *Science* 246:1275; and Orlandi et al. 1989 *PNAS* 86:3833). After immunizing an animal with an immunogen as described above, the antibody repertoire of the resulting B-cell pool is cloned. Methods are generally known for obtaining the DNA sequence of the variable regions of a diverse population of immunoglobulin molecules by using a mixture of oligomer primers and PCR. For instance, mixed oligonucleotide primers corresponding to the 5' leader (signal peptide) sequences and/or framework 1 (FR1) sequences, as well as primer to a conserved 3' constant region primer can be used for PCR amplification of the heavy and light chain variable regions from a number of murine antibodies (Larrick et al., 1991, *Biotechniques* 11:152-156). A similar strategy can also be used to amplify human heavy and light chain variable regions from human antibodies (Larrick et al., 1991, *Methods: Companion to Methods in Enzymology* 2:106-110).

Chimeric antibodies, including chimeric immunoglobulin chains, can be produced by recombinant DNA techniques known in the art. For example, a gene

encoding the Fc constant region of a murine (or other species) monoclonal antibody molecule is digested with restriction enzymes to remove the region encoding the murine Fc; and the equivalent portion of a gene encoding a human Fc constant region is substituted (see Robinson et al., International Patent Publication PCT/US86/02269; Akira, et al., European Patent Application 184,187; Taniguchi, M., European Patent Application 171,496; Morrison et al., European Patent Application 173,494; Neuberger et al., International Application WO 86/01533; Cabilly et al. U.S. Patent No. 4,816,567; Cabilly et al., European Patent Application 125,023; Better et al. (1988 *Science* 240:1041-1043); Liu et al. (1987) *PNAS* 84:3439-3443; Liu et al., 1987, *J. Immunol.* 139:3521-3526; Sun et al. (1987) *PNAS* 84:214-218; Nishimura et al., 1987, *Canc. Res.* 47:999-1005; Wood et al. (1985) *Nature* 314:446-449; and Shaw et al., 1988, *J. Natl Cancer Inst.* 80:1553-1559).

An antibody or an immunoglobulin chain can be humanized by methods known in the art. Humanized antibodies, including humanized immunoglobulin chains, can be generated by replacing sequences of the Fv variable region which are not directly involved in antigen binding with equivalent sequences from human Fv variable regions. General methods for generating humanized antibodies are provided by Morrison, S. L., 1985, *Science* 229:1202-1207, by Oi et al., 1986, *BioTechniques* 4:214, and by Queen et al. US 5,585,089, US 5,693,761 and US 5,693,762, the contents of all of which are hereby incorporated by reference. Those methods include isolating, manipulating, and expressing the nucleic acid sequences that encode all or part of immunoglobulin Fv variable regions from at least one of a heavy or light chain. Sources of such nucleic acid are well known to those skilled in the art and, for example, may be obtained from a hybridoma producing an antibody against a predetermined target. The recombinant DNA encoding the humanized antibody, or fragment thereof, can then be cloned into an appropriate expression vector.

Humanized or CDR-grafted antibody molecules or immunoglobulins can be produced by CDR-grafting or CDR substitution, wherein one, two, or all CDR's of an immunoglobulin chain can be replaced. See e.g., U.S. Patent 5,225,539; Jones et al. 1986 *Nature* 321:552-525; Verhoeyan et al. 1988 *Science* 239:1534; Beidler et al. 1988 *J. Immunol.* 141:4053-4060; Winter US 5,225,539, the contents of all of which are hereby expressly incorporated by reference. Winter describes a CDR-grafting method

which may be used to prepare the humanized antibodies of the present invention (UK Patent Application GB 2188638A, filed on March 26, 1987; Winter US 5,225,539), the contents of which is expressly incorporated by reference. All of the CDR's of a particular human antibody may be replaced with at least a portion of a non-human CDR or only
5 some of the CDR's may be replaced with non-human CDR's. It is only necessary to replace the number of CDR's required for binding of the humanized antibody to a predetermined antigen.

Monoclonal, chimeric and humanized antibodies, which have been modified by, e.g., deleting, adding, or substituting other portions of the antibody, e.g., the constant
10 region, are also within the scope of the invention. For example, an antibody can be modified as follows: (i) by deleting the constant region; (ii) by replacing the constant region with another constant region, e.g., a constant region meant to increase half-life, stability or affinity of the antibody, or a constant region from another species or antibody class; or (iii) by modifying one or more amino acids in the constant region to alter, for
15 example, the number of glycosylation sites, effector cell function, Fc receptor (FcR) binding, complement fixation, among others.

Methods for altering an antibody constant region are known in the art. Antibodies with altered function, e.g. altered affinity for an effector ligand, such as FcR on a cell, or the C1 component of complement can be produced by replacing at least one
20 amino acid residue in the constant portion of the antibody with a different residue (see e.g., EP 388,151 A1, US 5,624,821 and US 5,648,260, the contents of all of which are hereby incorporated by reference). Similar type of alterations could be described which if applied to the murine, or other species immunoglobulin would reduce or eliminate these functions.

For example, it is possible to alter the affinity of an Fc region of an antibody (e.g., an IgG, such as a human IgG) for an FcR (e.g., Fc gamma R1), or for C1q binding by replacing the specified residue(s) with a residue(s) having an appropriate functionality on its side chain, or by introducing a charged functional group, such as glutamate or aspartate, or perhaps an aromatic non-polar residue such as phenylalanine, tyrosine,
25 tryptophan or alanine (see e.g., US 5,624,821).
30

Amino acid sequences of IL-21 polypeptides are publicly known. For example, the nucleotide sequence and amino acid sequence of a human IL-21 is available at

Genbank Acc. No. X_011082. The disclosed human IL-21 nucleotide sequence is presented below:

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      1 gctgaagtga aaacgagacc aaggtctagc tctactgttg gtacttatga
      gatccagtc
5      61 tggcaacatg gagaggattg tcatctgtct gatggtcac ttcttgggga
      cactggtcca
      121 caaatcaagc tcccaaggtc aagatcgcca catgattaga atgctcaac
      ttatagatat
      181 tgttgatcag ctgaaaaatt atgtgaatga ctgggtccct gaatttctgc
10     cagctccaga
      241 agatgtagag acaaactgtg agtggtcagc tttttcctgc tttcagaagg
      cccaactaaa
      301 gtcagcaa at acaggaaaca atgaaaggat aatcaatgta tcaattaaaa
      agctgaagag
15     361 gaaaccacct tccacaaatg cagggagaag acagaaacac agactaacat
      gcccttcagt
      421 tgattcttat gagaaaaaac caccacaaaga attcctagaa agattcaa at
      cacttctcca
      481 aaagatgatt catcagcatc tgtcctctag aacacacgga agtgaagatt
20     cctgaggatc
      541 taacttgcag ttggacacta tgttacatac totaatatag tagtgaaagt
      catttcttg
      601 tattccaagt ggaggag (SEQ ID NO:18)

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25 The amino acid sequence of the disclosed human IL-21 polypeptide is presented below:

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MRSSPGNMERIVICLMVIFLGTLVHKSSSQGQDRHMIRMRLIDIVDQLKNYVNDLVE
FLPAPEDVETNCEWSAFSCFQKAQLKSANTGNNERIINVS IKKLKRKPPSTNAGRRQKH
30 RLTCPSCDSEYKKPPKEFLERFKSLLOKMIHQHLSSRTHGSEDS (SEQ ID NO:19)

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The invention also encompasses nucleic acids that hybridize to the nucleotide sequence set forth in SEQ ID NO:1, SEQ ID NO:22, SEQ ID NO:24, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:30, SEQ ID NO:32, SEQ ID NO:34, SEQ ID NO:36, or SEQ ID NO:38, under highly stringent conditions (for example, 0.1X SSC at 65° C.).

35 Isolated polynucleotides which encode MU-1 proteins or fusion proteins, but which differ from the nucleotide sequence set forth in SEQ ID NO:1, SEQ ID NO:22, SEQ ID NO:24, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:30, SEQ ID NO:32, SEQ ID NO:34, SEQ ID NO:36, or SEQ ID NO:38, by virtue of the degeneracy of the genetic code are also encompassed by the present invention. Variations in the nucleotide
40 sequence as set forth in SEQ ID NO:1, SEQ ID NO:22, SEQ ID NO:24, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:30, SEQ ID NO:32, SEQ ID NO:34, SEQ ID NO:36, or SEQ ID NO:38, which are caused by point mutations or by induced modifications are also included in the invention.

The isolated polynucleotides of the invention may be operably linked to an expression control sequence such as the pMT2 or pED expression vectors disclosed in Kaufman et al., Nucleic Acids Res. 19, 4485-4490 (1991), in order to produce the MU-1 protein recombinantly. Many suitable expression control sequences are known in the art.

5 General methods of expressing recombinant proteins are also known and are exemplified in R. Kaufman, Methods in Enzymology 185, 537-566 (1990). As defined herein "operably linked" means enzymatically or chemically ligated to form a covalent bond between the isolated polynucleotide of the invention and the expression control sequence, in such a way that the MU-1 protein is expressed by a host cell which has been

10 transformed (transfected) with the ligated polynucleotide/expression control sequence.

The term "vector", as used herein, is intended to refer to a nucleic acid molecule capable of transporting another nucleic acid to which it has been linked. One type of vector is a "plasmid", which refers to a circular double stranded DNA loop into which additional DNA segments may be ligated. Another type of vector is a viral vector,

15 wherein additional DNA segments may be ligated into the viral genome. Certain vectors are capable of autonomous replication in a host cell into which they are introduced (*e.g.*, bacterial vectors having a bacterial origin of replication and episomal mammalian vectors). Other vectors (*e.g.*, non-episomal mammalian vectors) can be integrated into the genome of a host cell upon introduction into the host cell, and thereby are replicated

20 along with the host genome. Moreover, certain vectors are capable of directing the expression of genes to which they are operatively linked. Such vectors are referred to herein as "recombinant expression vectors" (or simply, "expression vectors"). In general, expression vectors of utility in recombinant DNA techniques are often in the form of plasmids. In the present specification, "plasmid" and "vector" may be used

25 interchangeably as the plasmid is the most commonly used form of vector. However, the invention is intended to include such other forms of expression vectors, such as viral vectors (*e.g.*, replication defective retroviruses, adenoviruses and adeno-associated viruses), which serve equivalent functions.

The term "regulatory sequence" is intended to include promoters, enhancers and

30 other expression control elements (*e.g.*, polyadenylation signals) that control the transcription or translation of the antibody chain genes. Such regulatory sequences are described, for example, in Goeddel; *Gene Expression Technology: Methods in*

Enzymology 185, Academic Press, San Diego, CA (1990). It will be appreciated by those skilled in the art that the design of the expression vector, including the selection of regulatory sequences may depend on such factors as the choice of the host cell to be transformed, the level of expression of protein desired, etc. Preferred regulatory sequences for mammalian host cell expression include viral elements that direct high levels of protein expression in mammalian cells, such as promoters and/or enhancers derived from FF-1a promoter and BGH poly A, cytomegalovirus (CMV) (such as the CMV promoter/enhancer), Simian Virus 40 (SV40) (such as the SV40 promoter/enhancer), adenovirus, (e.g., the adenovirus major late promoter (AdMLP)) and polyoma. For further description of viral regulatory elements, and sequences thereof, see e.g., U.S. Patent No. 5,168,062 by Stinski, U.S. Patent No. 4,510,245 by Bell *et al.* and U.S. Patent No. 4,968,615 by Schaffner *et al.*

The recombinant expression vectors of the invention may carry additional sequences, such as sequences that regulate replication of the vector in host cells (e.g., origins of replication) and selectable marker genes. The selectable marker gene facilitates selection of host cells into which the vector has been introduced (see e.g., U.S. Patents Nos. 4,399,216, 4,634,665 and 5,179,017, all by Axel *et al.*). For example, typically the selectable marker gene confers resistance to drugs, such as G418, hygromycin or methotrexate, on a host cell into which the vector has been introduced. Preferred selectable marker genes include the dihydrofolate reductase (DHFR) gene (for use in dhfr⁻ host cells with methotrexate selection/amplification) and the *neo* gene (for G418 selection).

A number of types of cells may act as suitable host cells for expression of the MU-1 protein or fusion protein thereof. Any cell type capable of expressing functional MU-1 protein may be used. Suitable mammalian host cells include, for example, monkey COS cells, Chinese Hamster Ovary (CHO) cells, human kidney 293 cells, human epidermal A431 cells, human Colo205 cells, 3T3 cells, CV-1 cells, other transformed primate cell lines, normal diploid cells, cell strains derived from in vitro culture of primary tissue, primary explants, HeLa cells, mouse L cells, BHK, HL-60, U937, HaK, Rat2, BaF3, 32D, FDCP-1, PC12, M1x or C2C12 cells.

The MU-1 protein or fusion protein thereof may also be produced by operably linking the isolated polynucleotide of the invention to suitable control sequences in one

or more insect expression vectors, and employing an insect expression system. Materials and methods for baculovirus/insect cell expression systems are commercially available in kit form from, e.g., Invitrogen, San Diego, Calif. U.S.A. (the MaxBac® kit), and such methods are well known in the art, as described in Summers and Smith, Texas
5 Agricultural Experiment Station Bulletin No. 1555 (1987), incorporated herein by reference. Soluble forms of the MU-1 protein may also be produced in insect cells using appropriate isolated polynucleotides as described above.

Alternatively, the MU-1 protein or fusion protein thereof may be produced in lower eukaryotes such as yeast or in prokaryotes such as bacteria. Suitable yeast strains
10 include *Saccharomyces cerevisiae*, *Schizosaccharomyces pombe*, *Kluyveromyces* strains, *Candida*, or any yeast strain capable of expressing heterologous proteins. Suitable bacterial strains include *Escherichia coli*, *Bacillus subtilis*, *Salmonella typhimurium*, or any bacterial strain capable of expressing heterologous proteins.

Expression in bacteria may result in formation of inclusion bodies incorporating
15 the recombinant protein. Thus, refolding of the recombinant protein may be required in order to produce active or more active material. Several methods for obtaining correctly folded heterologous proteins from bacterial inclusion bodies are known in the art. These methods generally involve solubilizing the protein from the inclusion bodies, then denaturing the protein completely using a chaotropic agent. When cysteine residues are
20 present in the primary amino acid sequence of the protein, it is often necessary to accomplish the refolding in an environment which allows correct formation of disulfide bonds (a redox system). General methods of refolding are disclosed in Kohno, Meth. Enzym., 185:187-195 (1990). EP 0433225 and copending application U.S. Ser. No. 08/163,877 describe other appropriate methods.

25 The MU-1 protein or fusion protein thereof may also be expressed as a product of transgenic animals, e.g., as a component of the milk of transgenic cows, goats, pigs, or sheep which are characterized by somatic or germ cells containing a polynucleotide sequence encoding the MU-1 protein or fusion protein thereof.

The MU-1 protein or fusion protein thereof may be prepared by growing a culture
30 transformed host cells under culture conditions necessary to express the desired protein. The resulting expressed protein may then be purified from the culture medium or cell extracts. Soluble forms of the MU-1 protein or fusion protein thereof can be purified

from conditioned media. Membrane-bound forms of MU-1 protein of the invention can be purified by preparing a total membrane fraction from the expressing cell and extracting the membranes with a non-ionic detergent such as Triton X-100.

The MU-1 protein or fusion protein can be purified using methods known to those skilled in the art. For example, the MU-1 protein of the invention can be concentrated using a commercially available protein concentration filter, for example, an Amicon or Millipore Pellicon ultrafiltration unit. Following the concentration step, the concentrate can be applied to a purification matrix such as a gel filtration medium. Alternatively, an anion exchange resin can be employed, for example, a matrix or substrate having pendant diethylaminoethyl (DEAE) or polyethyleimine (PEI) groups. The matrices can be acrylamide, agarose, dextran, cellulose or other types commonly employed in protein purification. Alternatively, a cation exchange step can be employed. Suitable cation exchangers include various insoluble matrices comprising sulfopropyl or carboxymethyl groups. Sulfopropyl groups are preferred (e.g., S-Sepharose® columns). The purification of the MU-1 protein or fusion protein from culture supernatant may also include one or more column steps over such affinity resins as concanavalin A-agarose, heparin-toyopearl® or Cibacrom blue 3GA Sepharose®; or by hydrophobic interaction chromatography using such resins as phenyl ether, butyl ether, or propyl ether; or by immunoaffinity chromatography. Finally, one or more reverse-phase high performance liquid chromatography (RP-HPLC) steps employing hydrophobic RP-HPLC media, e.g., silica gel having pendant methyl or other aliphatic groups, can be employed to further purify the MU-1 protein. Affinity columns including antibodies to the MU-1 protein can also be used in purification in accordance with known methods. Some or all of the foregoing purification steps, in various combinations or with other known methods, can also be employed to provide a substantially purified isolated recombinant protein. Preferably, the isolated MU-1 protein is purified so that it is substantially free of other mammalian proteins.

MU-1 proteins or fusion proteins of the invention may also be used to screen for agents which are capable of binding to MU-1. Binding assays using a desired binding protein, immobilized or not, are well known in the art and may be used for this purpose using the MU-1 protein of the invention. Purified cell based or protein based (cell free) screening assays may be used to identify such agents. For example, MU-1 protein may

be immobilized in purified form on a carrier and binding or potential ligands to purified MU-1 protein may be measured.

Pharmaceutical Compositions

IL-21/IL-21R-agonists or antagonists may be used as a pharmaceutical composition when combined with a pharmaceutically acceptable carrier. Such a composition may contain, in addition to the IL-21/IL-21R-agonists or antagonists and carrier, various diluents, fillers, salts, buffers, stabilizers, solubilizers, and other materials well known in the art. The term "pharmaceutically acceptable" means a non-toxic material that does not interfere with the effectiveness of the biological activity of the active ingredient(s). The characteristics of the carrier will depend on the route of administration.

The pharmaceutical composition of the invention may also contain cytokines, lymphokines, or other hematopoietic factors such as M-CSF, GM-CSF, IL-1, IL-2, IL-3, IL-4, IL-5, IL-6, IL-7, IL-8, IL-9, IL-10, IL-11, IL-12, IL-14, IL-15, G-CSF, stem cell factor, and erythropoietin. The pharmaceutical composition may also include anti-cytokine antibodies as described in more detail below. The pharmaceutical composition may contain thrombolytic or anti-thrombotic factors such as plasminogen activator and Factor VIII. The pharmaceutical composition may further contain other anti-inflammatory agents as described in more detail below. Such additional factors and/or agents may be included in the pharmaceutical composition to produce a synergistic effect with an IL-21/IL-21R-agonists or antagonists, or to minimize side effects caused by the IL-21/IL-21R-agonists or antagonists. Conversely IL-21/IL-21R-agonists or antagonists may be included in formulations of the particular cytokine, lymphokine, other hematopoietic factor, thrombolytic or anti-thrombotic factor, or anti-inflammatory agent to minimize side effects of the cytokine, lymphokine, other hematopoietic factor, thrombolytic or anti-thrombotic factor, or anti-inflammatory agent.

The pharmaceutical composition of the invention may be in the form of a liposome in which IL-21/IL-21R-agonists or antagonists is combined, in addition to other pharmaceutically acceptable carriers, with amphipathic agents such as lipids which exist in aggregated form as micelles, insoluble monolayers, liquid crystals, or lamellar layers which in aqueous solution. Suitable lipids for liposomal formulation include, without limitation, monoglycerides, diglycerides, sulfatides, lysolecithin, phospholipids, saponin,

bile acids, and the like. Preparation of such liposomal formulations is within the level of skill in the art, as disclosed, for example, in U.S. Pat. No. 4,235,871; U.S. Pat. No. 4,501,728; U.S. Pat. No. 4,837,028; and U.S. Pat. No. 4,737,323, all of which are incorporated herein by reference.

- 5 As used herein, the term "therapeutically effective amount" means the total amount of each active component of the pharmaceutical composition or method that is sufficient to show a meaningful patient benefit, e.g., amelioration of symptoms of, healing of, or increase in rate of healing of such conditions. When applied to an individual active ingredient, administered alone, the term refers to that ingredient alone.
- 10 When applied to a combination, the term refers to combined amounts of the active ingredients that result in the therapeutic effect, whether administered in combination, serially or simultaneously.

- In practicing the method of treatment or use of the present invention, a therapeutically effective amount of an IL-21/IL-21R-agonist or antagonist is administered
- 15 to a subject, e.g., mammal (e.g., a human). An IL-21/IL-21R-agonists or antagonists may be administered in accordance with the method of the invention either alone or in combination with other therapies such as treatments employing cytokines, lymphokines or other hematopoietic factors, or anti-inflammatory agents. When co-administered with one or more agents, an IL-21- and/or IL-21R-binding agent may be administered either
- 20 simultaneously with the second agent, or sequentially. If administered sequentially, the attending physician will decide on the appropriate sequence of administering an IL-21/IL-21R-agonist or antagonist in combination with other agents.

- Administration of an IL-21/IL-21R-agonist or antagonist used in the pharmaceutical composition or to practice the method of the present invention can be
- 25 carried out in a variety of conventional ways, such as oral ingestion, inhalation, or cutaneous, subcutaneous, or intravenous injection. Intravenous administration to the patient is preferred.

- When a therapeutically effective amount of an IL-21/IL-21R-agonist or antagonist is administered orally, the binding agent will be in the form of a tablet, capsule, powder,
- 30 solution or elixir. When administered in tablet form, the pharmaceutical composition of the invention may additionally contain a solid carrier such as a gelatin or an adjuvant. The tablet, capsule, and powder contain from about 5 to 95% binding agent, and

preferably from about 25 to 90% binding agent. When administered in liquid form, a liquid carrier such as water, petroleum, oils of animal or plant origin such as peanut oil, mineral oil, soybean oil, or sesame oil, or synthetic oils may be added. The liquid form of the pharmaceutical composition may further contain physiological saline solution, dextrose or other saccharide solution, or glycols such as ethylene glycol, propylene glycol or polyethylene glycol. When administered in liquid form, the pharmaceutical composition contains from about 0.5 to 90% by weight of the binding agent, and preferably from about 1 to 50% the binding agent.

When a therapeutically effective amount of an IL-21/IL-21R-agonist or antagonist is administered by intravenous, cutaneous or subcutaneous injection, binding agent will be in the form of a pyrogen-free, parenterally acceptable aqueous solution. The preparation of such parenterally acceptable protein solutions, having due regard to pH, isotonicity, stability, and the like, is within the skill in the art. A preferred pharmaceutical composition for intravenous, cutaneous, or subcutaneous injection should contain, in addition to binding agent an isotonic vehicle such as Sodium Chloride Injection, Ringer's Injection, Dextrose Injection, Dextrose and Sodium Chloride Injection, Lactated Ringer's Injection, or other vehicle as known in the art. The pharmaceutical composition of the present invention may also contain stabilizers, preservatives, buffers, antioxidants, or other additive known to those of skill in the art.

The amount of an IL-21/IL-21R-agonist or antagonist in the pharmaceutical composition of the present invention will depend upon the nature and severity of the condition being treated, and on the nature of prior treatments that the patient has undergone. Ultimately, the attending physician will decide the amount of binding agent with which to treat each individual patient. Initially, the attending physician will administer low doses of binding agent and observe the patient's response. Larger doses of binding agent may be administered until the optimal therapeutic effect is obtained for the patient, and at that point the dosage is not generally increased further. It is contemplated that the various pharmaceutical compositions used to practice the method of the present invention should contain about 0.1 μ g to about 100 mg IL-21/IL-21R-agonist or antagonist per kg body weight.

The duration of intravenous therapy using the pharmaceutical composition of the present invention will vary, depending on the severity of the disease being treated and the

condition and potential idiosyncratic response of each individual patient. It is contemplated that the duration of each application of the IL-21/IL-21R-agonist or antagonist will be in the range of 12 to 24 hours of continuous intravenous administration. Ultimately the attending physician will decide on the appropriate
5 duration of intravenous therapy using the pharmaceutical composition of the present invention.

The polynucleotide and proteins of the present invention are expected to exhibit one or more of the uses or biological activities (including those associated with assays cited herein) identified below. Uses or activities described for proteins of the present
10 invention may be provided by administration or use of such proteins or by administration or use of polynucleotides encoding such proteins (such as, for example, in gene therapies or vectors suitable for introduction of DNA).

Uses of IL-21/IL-21R-Agonists to Enhance an Immune Response

In one aspect, the present invention provides methods for increasing immune cell,
15 e.g., T-cell (e.g., CD8+ T cell) proliferation by contacting an immune cell or a population of immune cells with an IL-21 or IL-21R binding agent which potentiates or enhances the activity of an IL-21 polypeptide. These methods are based, at least in part, on the finding that an agonist IL-21R binding agent stimulated, primarily via stimulation of CD8+T cells, innate and adaptive immunity *in vivo* against immunogenic and non-immunogenic
20 tumor cells (Example 9). The methods are also based, in part, on the finding that IL-21 induces proliferation of antigen- or anti-CD3 antibody stimulated thymocytes, lymph node T cells, CD4+ T cells, or CD8+ T cells. Applicants also show that proliferation of T cells to alloantigens can be increased in the presence of IL-21. In addition, IL-21 may also enhance proliferation and/or differentiation of CD8+ T cells (Example 10). For
25 example, priming of CD8+ T cells in the presence of IL-21 can generate effector cells with enhanced lytic (CTL) activity and/or increased ability to secrete cytokines, e.g., IFN γ . IL-21- or an IL-21R binding agent that stimulates IL-21/IL-21R activity can be used to induce the proliferation and/or cytokine secretion of macrophages. IL-21 also has effects on memory T cells and antigen presenting cells (APCs). IL-21 has also been
30 found to be produced by activated CD4⁺ cells. Accordingly, binding agents that stimulate the IL-21/IL-21R pathway can be used by themselves or in combination with an antigen,

e.g., as an adjuvant (e.g., a vaccine adjuvant), to up-regulate an immune response *in vivo*, e.g., for example, for use in treating cancer and infectious disorders.

In one embodiment, agonistic IL-21/IL-21R agonists may be useful in the treatment of various immune deficiencies and disorders (including severe combined immunodeficiency (SCID)), e.g., in regulating (up or down) growth and proliferation of T and/or B lymphocytes, as well as effecting the cytolytic activity of NK cells and other cell populations. These immune deficiencies may be genetic or be caused by viral (e.g., HIV) as well as bacterial or fungal infections, or may result from autoimmune disorders. More specifically, infectious diseases caused by viral, bacterial, fungal or other infection may be treatable using a protein of the present invention, including infections by HIV, hepatitis viruses, herpesviruses, mycobacteria, *Leishmania* spp., malaria spp. and various fungal infections such as candidiasis. Of course, in this regard, a protein of the present invention may also be useful where a boost to the immune system generally may be desirable, i.e., in the treatment of cancer.

Upregulation of an antigen function (preferably a B lymphocyte antigen function), as a means of up regulating immune responses, may also be useful in therapy. Upregulation of immune responses may be in the form of enhancing an existing immune response or eliciting an initial immune response. For example, enhancing an immune response through stimulating B lymphocyte antigen function may be useful in cases of viral infection. In addition, systemic viral diseases such as influenza, the common cold, and encephalitis might be alleviated by the administration of stimulatory forms of B lymphocyte antigens systemically.

Alternatively, anti-viral immune responses may be enhanced in an infected patient by removing T cells from the patient, costimulating the T cells *in vitro* with viral antigen-pulsed APCs either expressing a peptide of the present invention or together with a stimulatory form of a soluble peptide of the present invention and reintroducing the *in vitro* activated T cells into the patient. Another method of enhancing anti-viral immune responses would be to isolate infected cells from a patient, transfect them with a nucleic acid encoding a protein of the present invention as described herein such that the cells express all or a portion of the protein on their surface, and reintroduce the transfected cells into the patient. The infected cells would now be capable of delivering a costimulatory signal to, and thereby activate, T cells *in vivo*.

In another application, up regulation or enhancement of antigen function may be useful in the induction of tumor immunity. Tumor cells (e.g., sarcoma, melanoma, lymphoma, leukemia, neuroblastoma, carcinoma) transfected with a nucleic acid encoding at least one peptide of the present invention can be administered to a subject to overcome tumor-specific tolerance in the subject. If desired, the tumor cell can be transfected to express a combination of peptides. For example, tumor cells obtained from a patient can be transfected *ex vivo* with an expression vector directing the expression of an IL-21/IL-21R agonist, alone or in combination with, a peptide having B7-2-like activity alone, or in conjunction with a peptide having B7-1-like activity and/or B7-3-like activity. The transfected tumor cells are returned to the patient to result in expression of the peptides on the surface of the transfected cell. Alternatively, gene therapy techniques can be used to target a tumor cell for transfection *in vivo*.

The presence of an IL-21/IL-21R agonist, in combination with a peptide having the activity of a B lymphocyte antigen(s) on the surface of the tumor cell provides the necessary costimulation signal to T cells to induce a T cell mediated immune response against the transfected tumor cells. In addition, tumor cells which lack MHC class I or MHC class II molecules, or which fail to reexpress sufficient amounts of MHC class I or MHC class II molecules, can be transfected with nucleic acid encoding all or a portion of (e.g., a cytoplasmic-domain truncated portion) of an MHC class .alpha. a chain protein and β_2 microglobulin protein or an MHC class II α chain protein and an MHC class II β chain protein to thereby express MHC class I or MHC class II proteins on the cell surface. Expression of the appropriate class I or class II MHC in conjunction with an IL-21/IL-21R agonist, and/or a peptide having the activity of a B lymphocyte antigen (e.g., B7-1, B7-2, B7-3) induces a T cell mediated immune response against the transfected tumor cell. Optionally, a gene encoding an antisense construct which blocks expression of an MHC class II associated protein, such as the invariant chain, can also be cotransfected with a DNA encoding an IL-21/IL-21R agonist and/or a peptide having the activity of a B lymphocyte antigen to promote presentation of tumor associated antigens and induce tumor specific immunity. Thus, the induction of a T-cell mediated immune response in a human subject may be sufficient to overcome tumor-specific tolerance in the subject.

In other embodiments, the IL-21/IL-21R agonists can be used as vaccine adjuvants. Adjuvants are immune modulating compounds that have the ability to enhance and/or steer the development and profile of immune responses against various antigens that are themselves poorly immunogenic. Cytokines and/or lymphokines can be used as adjuvants. The appropriate selection of adjuvants can induce good humoral and cellular immune responses that would not develop in the absence of adjuvant. In particular, adjuvants have significant effects in enhancing the immune response to subunit and peptide antigens in vaccines. Their stimulatory activity is also beneficial to the development of antigen-specific immune responses directed against protein antigens. For a variety of antigens that require strong mucosal responses, high serum titers, induction of CTL and vigorous cellular responses, adjuvant and cytokine/lymphokine combinations provide stimuli that are not provided by most antigen preparations.

As used herein, the phrase "vaccine adjuvant" or "vaccine therapy" is intended to mean the use of an IL-21/IL-21R agonist, e.g., an IL-21 polypeptide or polynucleotide encoding the same, in combination with an antigen (e.g., viral, parasitic and bacterial polypeptides, proteins or peptides), or other antigens (e.g., tumor or cancer cell polypeptides, proteins or peptides) or polynucleotides encoding the antigen to enhance, suppress or otherwise modulate an immune response to the antigen. For the purpose of this definition, "combination" shall mean use in conjunction with, simultaneous with (combined or uncombined) or sequentially with an antigen.

The term "vaccine adjuvant composition" refers to a vaccine adjuvant that additionally includes immunologically acceptable diluents or carriers in a conventional manner to prepare injectable liquid solutions or suspensions. The vaccine adjuvant composition may additionally include agents that further enhance an immune response elicited by IL-21. For example, the vaccine adjuvant composition may additionally include 3-O-deacylated monophosphoryl lipid A (MPL™) or monophosphoryl lipid A and derivatives and analogs thereof. MPL™ can be used in a range of 1-100µg/dose.

The antigens used for vaccine therapy include proteins, peptides or polypeptides derived from immunogenic and non-immunogenic proteins, as well as any of the following: saccharides, proteins, poly- or oligonucleotides, or other macromolecular components, or fragments thereof. As used in this section, a "peptide" comprises a series of at least six amino acids and contains at least one antigenic determinant, while a

"polypeptide" is a longer molecule than a peptide, but does not constitute a full-length protein. As used herein, a "fragment" comprises a portion, but less than all of a saccharide, protein, poly- or oligonucleotide, or other macromolecular components.

As used herein, the term "effective adjuvanting amount" means a dose of the
5 combination of adjuvants described herein, which is suitable to elicit an increased immune response in a vertebrate host. The particular dosage will depend in part upon the age, weight and medical condition of the host, as well as on the method of administration and the antigen.

The vaccine adjuvant composition of the invention can be administered to a
10 human or non-human vertebrate by a variety of routes, including, but not limited to, intranasal, oral, vaginal, rectal, parenteral, intradermal, transdermal (see, e.g., International application WO 98/20734 (44), which is hereby incorporated by reference), intramuscular, intraperitoneal, subcutaneous, intravenous and intraarterial. The amount of the antigen component or components of the antigenic composition will vary
15 depending in part upon the identity of the antigen, as well as upon the age, weight and medical condition of the host, as well as on the method of administration. Again, suitable doses are readily determined by persons skilled in the art. It is preferable, although not required, that the antigen and the combination of adjuvants be administered at the same time. The number of doses and the dosage regimen for the antigenic
20 composition are also readily determined by persons skilled in the art. In some instances, the adjuvant properties of the combination of adjuvants may reduce the number of doses needed or the time course of the dosage regimen.

The combinations of adjuvants of this invention are suitable for use in combination with wide variety of antigens from a wide variety of pathogenic
25 microorganisms, including but not limited to those from viruses, bacteria, fungi or parasitic microorganisms that infect humans and non-human vertebrates, or from a cancer cell or tumor cell (e.g., sarcoma, melanoma, lymphoma, leukemia, neuroblastoma, carcinoma). The antigen may comprise peptides or polypeptides derived from proteins, as well as fragments of any of the following: saccharides, proteins, poly- or
30 oligonucleotides, cancer or tumor cells, allergens, amyloid peptide protein, or other macromolecular components. In some instances, more than one antigen is included in the antigenic composition.

Desirable viral vaccines containing the adjuvant combinations of this invention include those directed to the prevention and/or treatment of disease caused by, without limitation, Human immunodeficiency virus, Simian immunodeficiency virus, Respiratory syncytial virus, Parainfluenza virus types 1-3, Influenza virus, Herpes simplex virus, Human cytomegalovirus, Hepatitis A virus, Hepatitis B virus, Hepatitis C virus, Human papillomavirus, poliovirus, rotavirus, caliciviruses, Measles virus, Mumps virus, Rubella virus, adenovirus, rabies virus, canine distemper virus, rinderpest virus, coronavirus, parvovirus, infectious rhinotracheitis viruses, feline leukemia virus, feline infectious peritonitis virus, avian infectious bursal disease virus, Newcastle disease virus, Marek's disease virus, porcine respiratory and reproductive syndrome virus, equine arteritis virus and various Encephalitis viruses. In one embodiment, the IL-21/IL-21R agonists is administered in combination with a TNF antagonist, e.g., a TNF antagonist as described herein, to treat a Hepatitis C virus infection.

Desirable bacterial vaccines containing the adjuvant combinations of this invention include those directed to the prevention and/or treatment of disease caused by, without limitation, *Haemophilus influenzae* (both typable and nontypable), *Haemophilus somnus*, *Moraxella catarrhalis*, *Streptococcus pneumoniae*, *Streptococcus pyogenes*, *Streptococcus agalactiae*, *Streptococcus faecalis*, *Helicobacter pylori*, *Neisseria meningitidis*, *Neisseria gonorrhoeae*, *Chlamydia trachomatis*, *Chlamydia pneumoniae*, *Chlamydia psittaci*, *Bordetella pertussis*, *Salmonella typhi*, *Salmonella typhimurium*, *Salmonella choleraesuis*, *Escherichia coli*, *Shigella*, *Vibrio cholerae*, *Corynebacterium diphtheriae*, *Mycobacterium tuberculosis*, *Mycobacterium avium*-*Mycobacterium intracellulare* complex, *Proteus mirabilis*, *Proteus vulgaris*, *Staphylococcus aureus*, *Clostridium tetani*, *Leptospira interrogans*, *Borrelia burgdorferi*, *Pasteurella haemolytica*, *Pasteurella multocida*, *Actinobacillus pleuropneumoniae* and *Mycoplasma gallisepticum*.

Desirable vaccines against fungal pathogens containing the adjuvant combinations of this invention include those directed to the prevention and/or treatment of disease caused by, without limitation, *Aspergillus*, *Blastomyces*, *Candida*, *Coccidioides*, *Cryptococcus* and *Histoplasma*.

Desirable vaccines against parasites containing the adjuvant combinations of this invention include those directed to the prevention and/or treatment of disease caused by,

without limitation, *Leishmania major*, *Ascaris*, *Trichuris*, *Giardia*, *Schistosoma*, *Cryptosporidium*, *Trichomonas*, *Toxoplasma gondii* and *Pneumocystis carinii*.

Desirable vaccines for eliciting a therapeutic or prophylactic anti-cancer effect in a vertebrate host, which contain the adjuvant combinations of this invention, include
5 those utilizing a cancer antigen or tumor-associated antigen including, without limitation, prostate specific antigen (PSA), prostate-specific membrane antigen (PSMA), carcino-embryonic antigen (CEA), MUC-1, Her2, CA-125, MAGE-3, EGFR, HELP, GCC, CD66-c, prostasin, TMPRSS3, TADG 12 and TADG 15.

Desirable vaccines for moderating responses to allergens in a vertebrate host,
10 which contain the adjuvant combinations of this invention, include those containing an allergen or fragment thereof. Examples of such allergens are described in United States Patent Number 5,830,877 (45) and published International Patent Application Number WO 99/51259 (46), which are hereby incorporated by reference, and include pollen, insect venoms, animal dander, fungal spores and drugs (such as penicillin). The vaccines
15 interfere with the production of IgE antibodies, a known cause of allergic reactions.

Desirable vaccines for preventing or treating disease characterized by amyloid deposition in a vertebrate host, which contain the adjuvant combinations of this invention, include those containing portions of amyloid peptide protein (APP). This disease is referred to variously as Alzheimer's disease, amyloidosis or amyloidogenic
20 disease. Thus, the vaccines of this invention include the adjuvant combinations of this invention plus A β peptide, as well as fragments of A β peptide and antibodies to A β peptide or fragments thereof.

In the case of HIV and SIV, the antigenic compositions comprise at least one protein, polypeptide, peptide or fragment derived from said protein. In some instances,
25 multiple HIV or SIV proteins, polypeptides, peptides and/or fragments are included in the antigenic composition.

The adjuvant combination formulations of this invention are also suitable for inclusion as an adjuvant in polynucleotide vaccines (also known as DNA vaccines). Such vaccines may further include facilitating agents such as bupivacaine (see U.S.
30 Patent Number 5,593,972 (49), which is hereby incorporated by reference).

Uses of IL-21/IL-21R Antagonists to Decrease Immune Cell Activity

In yet another aspect, the invention features a method for inhibiting the activity of an immune cell, e.g., mature T cells (mature CD8+T cells, mature CD4+ T cells), mature NK cells, B cells, macrophages and megakaryocytes, or a population thereof, by contacting a population of T cells with an IL-21/IL-21R antagonist in an amount
5 sufficient to inhibit the activity of the immune cell or population. Antagonists of IL-21 and/or IL-21R (e.g., a fusion protein or a neutralizing antibody, as described herein) can also be administered to subjects for which inhibition of an immune response is desired. These conditions include, e.g., autoimmune disorders (e.g., arthritic disorders), or organ transplantation.

10 Applicants have shown that a reduction of IL-21R activity by using a neutralizing fusion protein that includes the extracellular domain of the IL-21R fused to an Fc immunoglobulin region ameliorates inflammatory symptoms in mouse collagen-induced arthritis (CIA) animal models (Example 7). Expression of IL-21R mRNA is upregulated in the paws of CIA mice (Example 8). Accordingly, IL-21R binding agents that
15 antagonize IL-21/IL-21R activity can be used to induce immune suppression *in vivo*, e.g., for treating or preventing immune cell-associated pathologies, including transplant rejection and autoimmune disorders.

The IL-21R DNA also maps to the chromosomal locus for Crohn's disease. As a result, binding agents of the present invention may be used to treat Crohn's disease and
20 other inflammatory bowel diseases.

The subject method can also be used to modulate (e.g., inhibit) the activity, e.g., proliferation, differentiation, survival) of an immune or hematopoietic cell (e.g., a cell of myeloid, lymphoid, erythroid lineages, or precursor cells thereof), and, thus, can be used to treat or prevent a variety of immune disorders. Non-limiting examples of the disorders
25 that can be treated or prevented include, but are not limited to, transplant rejection, autoimmune diseases (including, for example, diabetes mellitus, arthritis (including rheumatoid arthritis, juvenile rheumatoid arthritis, osteoarthritis, psoriatic arthritis), multiple sclerosis, encephalomyelitis, myasthenia gravis, systemic lupus erythematosus, autoimmune thyroiditis, dermatitis (including atopic dermatitis and eczematous
30 dermatitis), psoriasis, Sjögren's Syndrome, Crohn's disease, aphthous ulcer, iritis, conjunctivitis, keratoconjunctivitis, ulcerative colitis, spondyloarthropathy, ankylosing spondylitis, intrinsic asthma, allergic asthma, cutaneous lupus erythematosus,

scleroderma, vaginitis, proctitis, drug eruptions, leprosy reversal reactions, erythema nodosum leprosum, autoimmune uveitis, allergic encephalomyelitis, acute necrotizing hemorrhagic encephalopathy, idiopathic bilateral progressive sensorineural hearing loss, aplastic anemia, pure red cell anemia, idiopathic thrombocytopenia, polychondritis, Wegener's granulomatosis, chronic active hepatitis, Stevens-Johnson syndrome, idiopathic sprue, lichen planus, Graves' disease, sarcoidosis, primary biliary cirrhosis, uveitis posterior, and interstitial lung fibrosis), graft-versus-host disease, and allergy such as, atopic allergy. Preferred disorders that can be treated using the binding agents of the invention include arthritic disorders (e.g., rheumatoid arthritis, juvenile rheumatoid arthritis, osteoarthritis, psoriatic arthritis, and ankylosing spondylitis (preferably, rheumatoid arthritis)), multiple sclerosis, type I diabetes, lupus (SLE), IBD, Crohn's disease, asthma, vasculitis, allergy, scleroderma and psoriasis.

In another embodiment, IL-21/IL-21R antagonists, alone or in combination with, other therapeutic agents as described herein (e.g., TNF antagonists) can be used to treat multiple myeloma and related B lymphocytic malignancies (Brenne, A. *et al.* (2002) *Blood* Vol. 99(10):3756-3762).

Using the IL-21/IL-21R antagonists, it is possible to modulate immune responses, in a number of ways. Down regulation may be in the form of inhibiting or blocking an immune response already in progress or may involve preventing the induction of an immune response. The functions of activated T cells may be inhibited by suppressing T cell responses or by inducing specific tolerance in T cells, or both. Immunosuppression of T cell responses is generally an active, non-antigen-specific, process that requires continuous exposure of the T cells to the suppressive agent. Tolerance, which involves inducing non-responsiveness or anergy in T cells, is distinguishable from immunosuppression in that it is generally antigen-specific and persists after exposure to the tolerizing agent has ceased. Operationally, tolerance can be demonstrated by the lack of a T cell response upon reexposure to specific antigen in the absence of the tolerizing agent.

Down regulating or preventing one or more antigen functions (including without limitation B lymphocyte antigen functions, e.g., preventing high level lymphokine synthesis by activated T cells, will be useful in situations of tissue, skin and organ transplantation and in graft-versus-host disease (GVHD). For example, blockage of T cell

function should result in reduced tissue destruction in tissue transplantation. Typically, in tissue transplants, rejection of the transplant is initiated through its recognition as foreign by T cells, followed by an immune reaction that destroys the transplant. The administration of an IL-21/IL-21R antagonist, in combination with a molecule which inhibits or blocks interaction of a B7 lymphocyte antigen with its natural ligand(s) on immune cells (such as a soluble, monomeric form of a peptide having B7-2 activity alone or in conjunction with a monomeric form of a peptide having an activity of another B lymphocyte antigen (e.g., B7-1, B7-3) or blocking antibody), prior to transplantation can lead to the binding of the molecule to the natural ligand(s) on the immune cells without transmitting the corresponding costimulatory signal. Blocking B lymphocyte antigen function in this matter prevents cytokine synthesis by immune cells, such as T cells, and thus acts as an immunosuppressant. Moreover, the lack of costimulation may also be sufficient to anergize the T cells, thereby inducing tolerance in a subject. Induction of long-term tolerance by B lymphocyte antigen-blocking reagents may avoid the necessity of repeated administration of these blocking reagents. To achieve sufficient immunosuppression or tolerance in a subject, it may also be necessary to block the function of a combination of B lymphocyte antigens.

The efficacy of particular blocking reagents in preventing organ transplant rejection or GVHD can be assessed using animal models that are predictive of efficacy in humans. Examples of appropriate systems which can be used include allogeneic cardiac grafts in rats and xenogeneic pancreatic islet cell grafts in mice, both of which have been used to examine the immunosuppressive effects of CTLA4Ig fusion proteins in vivo as described in Lenschow et al., *Science* 257:789-792 (1992) and Turka et al., *Proc. Natl. Acad. Sci U.S.A.*, 89:11102-11105 (1992). In addition, murine models of GVHD (see Paul ed., *Fundamental Immunology*, Raven Press, New York, 1989, pp. 846-847) can be used to determine the effect of blocking B lymphocyte antigen function in vivo on the development of that disease.

Blocking antigen function may also be therapeutically useful for treating autoimmune diseases. Many autoimmune disorders are the result of inappropriate activation of T cells that are reactive against self tissue and which promote the production of cytokines and autoantibodies involved in the pathology of the diseases. Preventing the activation of autoreactive T cells may reduce or eliminate disease

symptoms. Administration of IL-21/IL-21R antagonists in combination with reagents which block costimulation of T cells by disrupting receptor:ligand interactions of B lymphocyte antigens can be used to inhibit T cell activation and prevent production of autoantibodies or T cell-derived cytokines which may be involved in the disease process.

- 5 Additionally, IL-21/IL-21R antagonists in combination with blocking reagents may induce antigen-specific tolerance of autoreactive T cells which could lead to long-term relief from the disease. The efficacy of these agents in preventing or alleviating autoimmune disorders can be determined using a number of well-characterized animal models of human autoimmune diseases. Examples include murine experimental
- 10 autoimmune encephalitis, systemic lupus erythematosus in MRL/lpr/lpr mice or NZB hybrid mice, murine autoimmune collagen arthritis, diabetes mellitus in NOD mice and BB rats, and murine experimental myasthenia gravis (see Paul ed., Fundamental Immunology, Raven Press, New York, 1989, pp. 840-856).

- In one embodiment, the IL-21/IL-21R antagonists, e.g., pharmaceutical
- 15 compositions thereof, are administered in combination therapy, i.e., combined with other agents, e.g., therapeutic agents, that are useful for treating pathological conditions or disorders, such as immune and inflammatory disorders. The term "in combination" in this context means that the agents are given substantially contemporaneously, either simultaneously or sequentially. If given sequentially, at the onset of administration of the
- 20 second compound, the first of the two compounds is preferably still detectable at effective concentrations at the site of treatment.

- For example, the combination therapy can include one or more IL-21/IL-21R antagonists, e.g., an antibody or an antigen-binding fragment thereof (e.g., a chimeric, humanized, human, or in vitro generated antibody or antigen-binding fragment thereof)
- 25 against IL-21 or IL-21 receptor, an IL-21 fusion protein, a soluble IL-21 receptor, peptide inhibitor or a small molecule inhibitor) co-formulated with, and/or co-administered with, one or more additional therapeutic agents, e.g., one or more cytokine and growth factor inhibitors, immunosuppressants, anti-inflammatory agents, metabolic inhibitors, enzyme inhibitors, and/or cytotoxic or cytostatic agents, as described in more detail below.

- 30 Furthermore, one or more IL-21/IL-21R antagonists described herein may be used in combination with two or more of the therapeutic agents described herein. Such combination therapies may advantageously utilize lower dosages of the administered

therapeutic agents, thus avoiding possible toxicities or complications associated with the various monotherapies. Moreover, the therapeutic agents disclosed herein act on pathways that differ from the IL-21/IL-21R receptor pathway, and thus are expected to enhance and/or synergize with the effects of the IL-21/IL-21R antagonists.

5 Preferred therapeutic agents used in combination with an IL-21/IL-21R antagonist are those agents that interfere at different stages in the autoimmune and subsequent inflammatory response. In one embodiment, one or more IL-21/IL-21R antagonist described herein may be co-formulated with, and/or co-administered with, one or more additional agents such as other cytokine or growth factor antagonists (e.g., soluble
10 receptors, peptide inhibitors, small molecules, ligand fusions); or antibodies or antigen-binding fragments thereof that bind to other targets (e.g., antibodies that bind to other cytokines or growth factors, their receptors, or other cell surface molecules); and anti-inflammatory cytokines or agonists thereof. Non-limiting examples of the agents that can be used in combination with the IL-21/IL-21R antagonists described herein, include,
15 but are not limited to, antagonists of one or more interleukins (ILs) or their receptors, e.g., antagonists of IL-1, IL-2, IL-6, IL-7, IL-8, IL-12, IL-13, IL-15, IL-16, IL-18, and IL-22; antagonists of cytokines or growth factors or their receptors, such as tumor necrosis factor (TNF), LT, EMAP-II, GM-CSF, FGF and PDGF. IL-21/IL-21R antagonists can also be combined with inhibitors of, e.g., antibodies to, cell surface molecules such as
20 CD2, CD3, CD4, CD8, CD25, CD28, CD30, CD40, CD45, CD69, CD80 (B7.1), CD86 (B7.2), CD90, or their ligands, including CD154 (gp39 or CD40L), or LFA-1/ICAM-1 and VLA-4/VCAM-1 (Yusuf-Makagiansar H. et al. (2002) *Med Res Rev* 22(2):146-67). Preferred antagonists that can be used in combination with IL-21/IL-21R antagonists described herein include antagonists of IL-1, IL-12, TNFa, IL-15, IL-17, IL-18, and IL-
25 22.

Examples of those agents include IL-12 antagonists, such as chimeric, humanized, human or in vitro generated antibodies (or antigen-binding fragments thereof) that bind to IL-12 (preferably human IL-12), e.g., the antibody disclosed in WO 00/56772, Genetics Institute/BASF); IL-12 receptor inhibitors, e.g., antibodies to human
30 IL-12 receptor; and soluble fragments of the IL-12 receptor, e.g., human IL-12 receptor. Examples of IL-15 antagonists include antibodies (or antigen-binding fragments thereof) against IL-15 or its receptor, e.g., chimeric, humanized, human or *in vitro* generated

antibodies to human IL-15 or its receptor, soluble fragments of the IL-15 receptor, and IL-15-binding proteins. Examples of IL-18 antagonists include antibodies, e.g., chimeric, humanized, human or in vitro generated antibodies (or antigen-binding fragments thereof), to human IL-18, soluble fragments of the IL-18 receptor, and IL-18 binding proteins (IL-18BP, Mallet et al. (2001) *Circ. Res.* 28). Examples of IL-1 antagonists include Interleukin-1-converting enzyme (ICE) inhibitors, such as Vx740, IL-1 antagonists, e.g., IL-1RA (ANIKINRA, AMGEN), sIL1RII (Immunex), and anti-IL-1 receptor antibodies (or antigen-binding fragments thereof).

Examples of TNF antagonists include chimeric, humanized, human or in vitro generated antibodies (or antigen-binding fragments thereof) to TNF (e.g., human TNF α), such as D2E7, (human TNF α antibody, U.S. 6,258,562; BASF), CDP-571/CDP-870/BAY-10-3356 (humanized anti-TNF α antibody; Celltech/Pharmacia), cA2 (chimeric anti-TNF α antibody; RemicadeTM, Centocor); anti-TNF antibody fragments (e.g., CPD870); soluble fragments of the TNF receptors, e.g., p55 or p75 human TNF receptors or derivatives thereof, e.g., 75 kDTNFR-IgG (75 kD TNF receptor-IgG fusion protein, EnbrelTM; Immunex; see e.g., *Arthritis & Rheumatism* (1994) Vol. 37, S295; *J. Invest. Med.* (1996) Vol. 44, 235A), p55 kDTNFR-IgG (55 kD TNF receptor-IgG fusion protein (Lenercept)); enzyme antagonists, e.g., TNF α converting enzyme (TACE) inhibitors (e.g., an alpha-sulfonyl hydroxamic acid derivative, WO 01/55112, and N-hydroxyformamide TACE inhibitor GW 3333, -005, or -022); and TNF-bp/s-TNFR (soluble TNF binding protein; see e.g., *Arthritis & Rheumatism* (1996) Vol. 39, No. 9 (supplement), S284; *Amer. J. Physiol. - Heart and Circulatory Physiology* (1995) Vol. 268, pp. 37-42). Preferred TNF antagonists are soluble fragments of the TNF receptors, e.g., p55 or p75 human TNF receptors or derivatives thereof, e.g., 75 kDTNFR-IgG, and TNF α converting enzyme (TACE) inhibitors.

In other embodiments, the IL-21-/IL21R binding agents described herein can be administered in combination with one or more of the following: IL-13 antagonists, e.g., soluble IL-13 receptors (sIL-13) and/or antibodies against IL-13; IL-2 antagonists, e.g., DAB 486-IL-2 and/or DAB 389-IL-2 (IL-2 fusion proteins; Seragen; see e.g., *Arthritis & Rheumatism* (1993) Vol. 36, 1223), and/or antibodies to IL-2R, e.g., anti-Tac (humanized anti-IL-2R; Protein Design Labs, *Cancer Res.* 1990 Mar 1;50(5):1495-502). Yet another combination includes IL-21 antagonists in combination with non-depleting

anti-CD4 inhibitors (IDEC-CE9.1/SB 210396 (non-depleting primatized anti-CD4 antibody; IDEC/SmithKline). Yet other preferred combinations include antagonists of the co-stimulatory pathway CD80 (B7.1) or CD86 (B7.2) including antibodies, soluble receptors or antagonistic ligands; as well as p-selectin glycoprotein ligand (PSGL), anti-inflammatory cytokines, e.g., IL-4 (DNAX/Schering); IL-10 (SCH 52000; recombinant IL-10 DNAX/Schering); IL-13 and TGF β , and agonists thereof (e.g., agonist antibodies).

In other embodiments, one or more IL-21-/IL21R binding agents can be co-formulated with, and/or co-administered with, one or more anti-inflammatory drugs, immunosuppressants, or metabolic or enzymatic inhibitors. Non-limiting examples of the drugs or inhibitors that can be used in combination with the IL-21 antagonists described herein, include, but are not limited to, one or more of: non-steroidal anti-inflammatory drug(s) (NSAIDs), e.g., ibuprofen, Tenidap (see e.g., *Arthritis & Rheumatism* (1996) Vol. 39, No. 9 (supplement), S280)), Naproxen (see e.g., *Neuro Report* (1996) Vol. 7, pp. 1209-1213), Meloxicam, Piroxicam, Diclofenac, and Indomethacin; Sulfasalazine (see e.g., *Arthritis & Rheumatism* (1996) Vol. 39, No. 9 (supplement), S281); corticosteroids such as prednisolone; cytokine suppressive anti-inflammatory drug(s) (CSAIDs); inhibitors of nucleotide biosynthesis, e.g., inhibitors of purine biosynthesis, folate antagonists (e.g., methotrexate (N-[4-[(2,4-diamino-6-pteridiny)methyl]methylamino]benzoyl]-L-glutamic acid); and inhibitors of pyrimidine biosynthesis, e.g., dihydroorotate dehydrogenase (DHODH) inhibitors (e.g., leflunomide (see e.g., *Arthritis & Rheumatism* (1996) Vol. 39, No. 9 (supplement), S131; *Inflammation Research* (1996) Vol. 45, pp. 103-107). Preferred therapeutic agents for use in combination with IL-21/IL-21R antagonists include NSAIDs, CSAIDs, (DHODH) inhibitors (e.g., leflunomide), and folate antagonists (e.g., methotrexate).

Examples of additional inhibitors include one or more of: corticosteroids (oral, inhaled and local injection); immunosuppressants, e.g., cyclosporin, tacrolimus (FK-506); and mTOR inhibitors, e.g., sirolimus (rapamycin) or rapamycin derivatives, e.g., soluble rapamycin derivatives (e.g., ester rapamycin derivatives, e.g., CCI-779 (Elit. L. (2002) *Current Opinion Investig. Drugs* 3(8):1249-53; Huang, S. et al.. (2002) *Current Opinion Investig. Drugs* 3(2):295-304); agents which interfere with signaling by proinflammatory cytokines such as TNF α or IL-1 (e.g. IRAK, NIK, IKK, p38 or MAP kinase inhibitors);

COX2 inhibitors, e.g., celecoxib and variants thereof, MK-966, see e.g., Arthritis & Rheumatism (1996) Vol. 39, No. 9 (supplement), S81); phosphodiesterase inhibitors, e.g., R973401 (phosphodiesterase Type IV inhibitor; see e.g., Arthritis & Rheumatism (1996) Vol. 39, No. 9 (supplement), S282)); phospholipase inhibitors, e.g., inhibitors of
5 cytosolic phospholipase 2 (cPLA2) (e.g., trifluoromethyl ketone analogs (U.S. 6,350,892)); inhibitors of vascular endothelial cell growth factor or growth factor receptor, e.g., VEGF inhibitor and/or VEGF-R inhibitor; and inhibitors of angiogenesis. Preferred therapeutic agents for use in combination with IL-21/IL-21R antagonists immunosuppressants, e.g., cyclosporin, tacrolimus (FK-506); and mTOR inhibitors, e.g.,
10 sirolimus (rapamycin) or rapamycin derivatives, e.g., soluble rapamycin derivatives (e.g., ester rapamycin derivatives, e.g., CCI-779; COX2 inhibitors, e.g., celecoxib and variants thereof; and phospholipase inhibitors, e.g., inhibitors of cytosolic phospholipase 2 (cPLA2) (e.g., trifluoromethyl ketone analogs)

Additional examples of therapeutic agents that can be combined with an IL-
15 21/IL-21R antagonist include one or more of: 6-mercaptopurines (6-MP); azathioprine sulphasalazine; mesalazine; olsalazine chloroquine/hydroxychloroquine; pencillamine; aurothiornalate (intramuscular and oral); azathioprine; cochlamine; beta-2 adrenoreceptor agonists (salbutamol, terbutaline, salmeteral); xanthines (theophylline, aminophylline); cromoglycate; nedocromil; ketotifen; ipratropium and oxitropium; mycophenolate
20 mofetil; adenosine agonists; antithrombotic agents; complement inhibitors; and adrenergic agents.

The use of the IL-21-/IL21R binding agents disclosed herein in combination with other therapeutic agents to treat or prevent specific immune disorders is discussed in further detail below.

25 Non-limiting examples of agents for treating or preventing arthritic disorders (e.g., rheumatoid arthritis, inflammatory arthritis, rheumatoid arthritis, juvenile rheumatoid arthritis, osteoarthritis and psoriatic arthritis), with which an IL-21-/IL21R binding agent can be combined include one or more of the following: IL-12 antagonists as described herein, NSAIDs; CSAIDs; TNF's, e.g., TNFa, antagonists as described
30 herein; non-depleting anti-CD4 antibodies as described herein; IL-2 antagonists as described herein; anti-inflammatory cytokines, e.g., IL-4, IL-10, IL-13 and TGfA, or agonists thereof; IL-1 or IL-1 receptor antagonists as described herein);

phosphodiesterase inhibitors as described herein; COX-2 inhibitors as described herein; Iloprost (see e.g., *Arthritis & Rheumatism* (1996) Vol. 39, No. 9 (supplement), S82); methotrexate; thalidomide (see e.g., *Arthritis & Rheumatism* (1996) Vol. 39, No. 9 (supplement), S282) and thalidomide-related drugs (e.g., Celgen); leflunomide; inhibitor of plasminogen activation, e.g., tranexamic acid; see e.g., *Arthritis & Rheumatism* (1996) Vol. 39, No. 9 (supplement), S284); cytokine inhibitor, e.g., T-614; see e.g., *Arthritis & Rheumatism* (1996) Vol. 39, No. 9 (supplement), S282); prostaglandin E1 (see e.g., *Arthritis & Rheumatism* (1996) Vol. 39, No. 9 (supplement), S282); azathioprine (see e.g., *Arthritis & Rheumatism* (1996) Vol. 39, No. 9 (supplement), S281); an inhibitor of interleukin-1 converting enzyme (ICE); zap-70 and/or lck inhibitor (inhibitor of the tyrosine kinase zap-70 or lck); an inhibitor of vascular endothelial cell growth factor or vascular endothelial cell growth factor receptor as described herein; an inhibitor of angiogenesis as described herein; corticosteroid anti-inflammatory drugs (e.g., SB203580); TNF-convertase inhibitors; interleukin-11 (see e.g., *Arthritis & Rheumatism* (1996) Vol. 39, No. 9 (supplement), S296); interleukin-13 (see e.g., *Arthritis & Rheumatism* (1996) Vol. 39, No. 9 (supplement), S308); interleukin-17 inhibitors (see e.g., *Arthritis & Rheumatism* (1996) Vol. 39, No. 9 (supplement), S120); gold; penicillamine; chloroquine; hydroxychloroquine; chlorambucil; cyclophosphamide; cyclosporine; total lymphoid irradiation; anti-thymocyte globulin; CD5-toxins; orally-administered peptides and collagen; lobenzarit disodium; Cytokine Regulating Agents (CRAs) HP228 and HP466 (Houghten Pharmaceuticals, Inc.); ICAM-1 antisense phosphorothioate oligodeoxynucleotides (ISIS 2302; Isis Pharmaceuticals, Inc.); soluble complement receptor 1 (TP10; T Cell Sciences, Inc.); prednisone; orgotein; glycosaminoglycan polysulphate; minocycline; anti-IL2R antibodies; marine and botanical lipids (fish and plant seed fatty acids; see e.g., DeLuca et al. (1995) *Rheum. Dis. Clin. North Am.* 21:759-777); auranofin; phenylbutazone; meclofenamic acid; flufenamic acid; intravenous immune globulin; zileuton; mycophenolic acid (RS-61443); tacrolimus (FK-506); sirolimus (rapamycin); amiprilose (therafectin); cladribine (2-chlorodeoxyadenosine); and azaribine. Preferred combinations include one or more IL-21 antagonists in combination with methotrexate or leflunomide, and in moderate or severe rheumatoid arthritis cases, cyclosporine.

Preferred examples of inhibitors to use in combination with UL-21/IL-21R to treat arthritic disorders include TNF antagonists (e.g., chimeric, humanized, human or *in vitro* generated antibodies, or antigen-binding fragments thereof, that bind to TNF; soluble fragments of a TNF receptor, e.g., p55 or p75 human TNF receptor or derivatives thereof, e.g., 75 kDTNFR-IgG (75 kD TNF receptor-IgG fusion protein, EnbrelTM), p55 kD TNF receptor-IgG fusion protein; TNF enzyme antagonists, e.g., TNFa converting enzyme (TACE) inhibitors); antagonists of IL-12, IL-15, IL-17, IL-18, IL-22; T cell and B cell depleting agents (e.g., anti-CD4 or anti-CD22 antibodies); small molecule inhibitors, e.g., methotrexate and leflunomide; sirolimus (rapamycin) and analogs thereof, e.g., CCI-779; Cox-2 and cPLA2 inhibitors; NSAIDs; p38 inhibitors, TPL-2, Mk-2 and NFkb inhibitors; RAGE or soluble RAGE; P-selectin or PSGL-1 inhibitors (e.g., small molecule inhibitors, antibodies thereto, e.g., antibodies to P-selectin); estrogen receptor beta (ERB) agonists or ERB-NFkb antagonists. Most preferred additional therapeutic agents that can be co-administered and/or co-formulated with one or more IL-21/IL-21R antagonists include one or more of: a soluble fragment of a TNF receptor, e.g., p55 or p75 human TNF receptor or derivatives thereof, e.g., 75 kDTNFR-IgG (75 kD TNF receptor-IgG fusion protein, EnbrelTM); methotrexate, leflunomide, or a sirolimus (rapamycin) or an analog thereof, e.g., CCI-779.

Non-limiting examples of agents for treating or preventing multiple sclerosis with which an IL-21-/IL21R binding agent can be combined include the following: interferons, e.g., interferon-alpha1a (e.g., AvonexTM; Biogen) and interferon-1b (BetaseronTM; Chiron/Berlex); Copolymer 1 (Cop-1; CopaxoneTM; Teva Pharmaceutical Industries, Inc.); hyperbaric oxygen; intravenous immunoglobulin; clabribine; TNF antagonists as described herein; corticosteroids; prednisolone; methylprednisolone; azathioprine; cyclophosphamide; cyclosporine; methotrexate; 4-aminopyridine; and tizanidine. Additional antagonists that can be used in combination with IL-21 include antibodies to or antagonists of other human cytokines or growth factors, for example, TNF, LT, IL-1, IL-2, IL-6, IL-7, IL-8, IL-12 IL-15, IL-16, IL-18, EMAP-11, GM-CSF, FGF, and PDGF. IL-21 antagonists as described herein can be combined with antibodies to cell surface molecules such as CD2, CD3, CD4, CD8, CD25, CD28, CD30, CD40, CD45, CD69, CD80, CD86, CD90 or their ligands. The IL-21 antagonists may also be combined with agents, such as methotrexate, cyclosporine, FK506, rapamycin,

mycophenolate mofetil, leflunomide, NSAIDs, for example, ibuprofen, corticosteroids such as prednisolone, phosphodiesterase inhibitors, adenosine agonists, antithrombotic agents, complement inhibitors, adrenergic agents, agents which interfere with signaling by proinflammatory cytokines as described herein, IL-1b converting enzyme inhibitors (e.g., Vx740), anti-P7s, PSGL, TACE inhibitors, T-cell signaling inhibitors such as kinase inhibitors, metal loproteinase inhibitors, sulfasalazine, azathioprine, 6-mercaptapurines, angiotensin converting enzyme inhibitors, soluble cytokine receptors and derivatives thereof, as described herein, and anti-inflammatory cytokines (e.g. IL-4, IL-10, IL-13 and TGF).

10 Preferred examples of therapeutic agents for multiple sclerosis with which the IL-21 antagonists can be combined include interferon-b, for example, IFNb-1a and IFNb-1b; copaxone, corticosteroids, IL-1 inhibitors, TNF inhibitors, antibodies to CD40 ligand and CD80, IL-12 antagonists.

Non-limiting examples of agents for treating or preventing inflammatory bowel disease or Crohn's disease with which an IL-21/IL-21R antagonist can be combined include the following: budesonide; epidermal growth factor; corticosteroids; cyclosporin, sulfasalazine; aminosalicylates; 6-mercaptapurine; azathioprine; metronidazole; lipooxygenase inhibitors; mesalamine; olsalazine; balsalazide; antioxidants; thromboxane inhibitors; IL-1 receptor antagonists; anti-IL-1 monoclonal antibodies; anti-IL-6 monoclonal antibodies; growth factors; elastase inhibitors; pyridinyl-imidazole compounds; TNF antagonists as described herein; IL-4, IL-10, IL-13 and/or TGFb cytokines or agonists thereof (e.g., agonist antibodies); interleukin-11; glucuronide- or dextran-conjugated prodrugs of prednisolone, dexamethasone or budesonide; ICAM-1 antisense phosphorothioate oligodeoxynucleotides (ISIS 2302; Isis Pharmaceuticals, Inc.); soluble complement receptor 1 (TP10; T Cell Sciences, Inc.); slow-release mesalazine; methotrexate; antagonists of Platelet Activating Factor (PAF); ciprofloxacin; and lignocaine.

In one embodiment, an IL-21/IL21R antagonists can be used in combination with one or more antibodies directed at other targets involved in regulating immune responses, e.g., transplant rejection or graft-v-host disease. Non-limiting examples of agents for treating or preventing immune responses with which an IL-21/IL21R antagonist of the invention can be combined include the following: antibodies against cell surface

molecules, including but not limited to CD25 (interleukin-2 receptor- α), CD11a (LFA-1), CD54 (ICAM-1), CD4, CD45, CD28/CTLA4, CD80 (B7-1) and/or CD86 (B7-2). In yet another embodiment, an IL-21/IL21R antagonist is used in combination with one or more general immunosuppressive agents, such as cyclosporin A or FK506.

5 Another aspect of the present invention accordingly relates to kits for carrying out the combined administration of the IL-21/IL21R antagonists with other therapeutic compounds. In one embodiment, the kit comprises one or more binding agents formulated in a pharmaceutical carrier, and at least one agent, e.g., therapeutic agent, formulated as appropriate, in one or more separate pharmaceutical preparations.

10 Assays for Measuring the Activity of IL-21/IL21R Agonists as Immune Activators

The activity of IL-21/IL21R agonists as activators of an immune system can, among other means, be measured by the following methods:

Suitable assays for thymocyte or splenocyte cytotoxicity include, without limitation, those described in: Current Protocols in Immunology, Ed by J. E. Coligan, A. M. Kruisbeek, D. H. Margulies, E. M. Shevach, W Strober, Pub. Greene Publishing Associates and Wiley-Interscience (Chapter 3, In Vitro assays for Mouse Lymphocyte Function 3.1-3.19; Chapter 7, Immunologic studies in Humans); Herrmann et al., Proc. Natl. Acad. Sci. U.S.A. 78:2488-2492, 1981; Herrmann et al., J. Immunol. 128:1968-1974, 1982; Handa et al., J. Immunol. 135:1564-1572, 1985; Takai et al., J. Immunol. 137:3494-3500, 1986; Takai et al., J. Immunol. 140:508-512, 1988; Herrmann et al., Proc. Natl. Acad. Sci. U.S.A. 78:2488-2492, 1981; Herrmann et al., J. Immunol. 128:1968-1974, 1982; Handa et al., J. Immunol. 135:1564-1572, 1985; Takai et al., J. Immunol. 137:3494-3500, 1986; Bowman et al., J. Virology 61:1992-1998; Takai et al., J. Immunol. 140:508-512, 1988; Bertagnoli et al., Cellular Immunology 133:327-341, 1991; Brown et al., J. Immunol. 153:3079-3092, 1994.

Assays for T-cell-dependent immunoglobulin responses and isotype switching (which will identify, among others, proteins that modulate T-cell dependent antibody responses and that affect Th1/Th2 profiles) include, without limitation, those described in: Maliszewski, J. Immunol. 144:3028-3033, 1990; and Assays for B cell function: In vitro antibody production, Mond, J. J. and Brunswick, M. In Current Protocols in Immunology. J. E. Coligan eds. Vol 1 pp. 3.8.1-3.8.16, John Wiley and Sons, Toronto. 1994.

Mixed lymphocyte reaction (MLR) assays (which will identify, among others, proteins that generate predominantly Th1 and CTL responses) include, without limitation, those described in: Current Protocols in Immunology, Ed by J. E. Coligan, A. M. Kruisbeek, D. H. Margulies, E. M. Shevach, W Strober, Pub. Greene Publishing Associates and Wiley-Interscience (Chapter 3, In Vitro assays for Mouse Lymphocyte Function 3.1-3.19; Chapter 7, Immunologic studies in Humans); Takai et al., J. Immunol. 137:3494-3500, 1986; Takai et al., J. Immunol. 140:508-512, 1988; Bertagnolli et al., J. Immunol. 149:3778-3783, 1992.

Dendritic cell-dependent assays (which will identify, among others, proteins expressed by dendritic cells that activate naive T-cells) include, without limitation, those described in: Guery et al., J. Immunol. 134:536-544, 1995; Inaba et al., Journal of Experimental Medicine 173:549-559, 1991; Macatonia et al., Journal of Immunology 154:5071-5079, 1995; Porgador et al., Journal of Experimental Medicine 182:255-260, 1995; Nair et al., Journal of Virology 67:4062-4069, 1993; Huang et al., Science 264:961-965, 1994; Macatonia et al., Journal of Experimental Medicine 169:1255-1264, 1989; Bhardwaj et al., Journal of Clinical Investigation 94:797-807, 1994; and Inaba et al., Journal of Experimental Medicine 172:631-640, 1990.

Assays for lymphocyte survival/apoptosis (which will identify, among others, proteins that prevent apoptosis after superantigen induction and proteins that regulate lymphocyte homeostasis) include, without limitation, those described in: Darzynkiewicz et al., Cytometry 13:795-808, 1992; Gorczyca et al., Leukemia 7:659-670, 1993; Gorczyca et al., Cancer Research 53:1945-1951, 1993; Itoh et al., Cell 66:233-243, 1991; Zacharchuk, Journal of Immunology 145:4037-4045, 1990; Zamai et al., Cytometry 14:891-897, 1993; Gorczyca et al., International Journal of Oncology 1:639-648, 1992.

Assays for proteins that influence early steps of T-cell commitment and development include, without limitation, those described in: Antica et al., Blood 84:111-117, 1994; Fine et al., Cellular Immunology 155:111-122, 1994; Galy et al., Blood 85:2770-2778, 1995; Toki et al., Proc. Nat. Acad Sci. U.S.A. 88:7548-7551, 1991.

Assays for Measuring the Activity of IL-21/IL21R Agonists or Antagonists as Modulators of Cytokine Production and Cell Proliferation/Differentiation

The activity of IL-21/IL21R agonists or antagonists as modulator of cytokine production and cell proliferation/differentiation can be tested using any one of a number

of routine factor dependent cell proliferation assays for cell lines including, without limitation, 32D, DA2, DA1G, T10, B9, B9/11, BaF3, MC9/G, M+(preB M+), 2E8, RB5, DA1, 123, T1165, HT2, CTLL2, TF-1, Mo7e and CMK.

- Assays for T-cell or thymocyte proliferation include without limitation those
- 5 described in: Current Protocols in Immunology, Ed by J. E. Coligan, A. M. Kruisbeek, D. H. Margulies, E. M. Shevach, W Strober, Pub. Greene Publishing Associates and Wiley-Interscience (Chapter 3, In Vitro assays for Mouse Lymphocyte Function 3.1-3.19; Chapter 7, Immunologic studies in Humans); Takai et al., J. Immunol. 137:3494-3500, 1986; Bertagnolli et al., J. Immunol. 145:1706-1712, 1990; Bertagnolli et al., Cellular
- 10 Immunology 133:327-341, 1991; Bertagnolli, et al., J. Immunol. 149:3778-3783, 1992; Bowman et al., J. Immunol. 152: 1756-1761, 1994.

- Assays for cytokine production and/or proliferation of spleen cells, lymph node cells or thymocytes include, without limitation, those described in: Polyclonal T cell stimulation, Kruisbeek, A. M. and Shevach, E. M. In Current Protocols in Immunology. J. E.e.a.
- 15 Coligan eds. Vol 1 pp. 3.12.1-3.12.14, John Wiley and Sons, Toronto. 1994; and Measurement of mouse and human Interferon .gamma., Schreiber, R. D. In Current Protocols in Immunology. J. E.e.a. Coligan eds. Vol 1 pp. 6.8.1-6.8.8, John Wiley and Sons, Toronto. 1994.

- Assays for proliferation and differentiation of hematopoietic and lymphopoietic
- 20 cells include, without limitation, those described in: Measurement of Human and Murine Interleukin 2 and Interleukin 4, Bottomly, K., Davis, L. S. and Lipsky, P. E. In Current Protocols in Immunology. J. E.e.a. Coligan eds. Vol 1 pp. 6.3.1-6.3.12, John Wiley and Sons, Toronto. 1991; deVries et al., J. Exp. Med. 173:1205-1211, 1991; Moreau et al., Nature 336:690-692, 1988; Greenberger et al., Proc. Natl. Acad. Sci. U.S.A. 80:2931-
- 25 2938, 1983; Measurement of mouse and human interleukin 6--Nordan, R. In Current Protocols in Immunology. J. E.e.a. Coligan eds. Vol 1 pp. 6.6.1-6.6.5, John Wiley and Sons, Toronto. 1991; Smith et al., Proc. Natl. Acad. Sci. U.S.A. 83:1857-1861, 1986; Measurement of human Interleukin 11--Bennett, F., Giannotti, J., Clark, S. C. and Turner, K. J. In Current Protocols in Immunology. J. E.e.a. Coligan eds. Vol 1 pp. 6.15.1
- 30 John Wiley and Sons, Toronto. 1991; Measurement of mouse and human Interleukin 9--Ciarletta, A., Giannotti, J., Clark, S. C. and Turner, K. J. In Current Protocols in

Immunology. J. E.e.a. Coligan eds. Vol 1 pp. 6.13.1, John Wiley and Sons, Toronto. 1991.

Assays for T-cell clone responses to antigens (which will identify, among others, proteins that affect APC-T cell interactions as well as direct T-cell effects by measuring proliferation and cytokine production) include, without limitation, those described in: Current Protocols in Immunology, Ed by J. E. Coligan, A. M. Kruisbeek, D. H. Margulies, E. M. Shevach, W Strober, Pub. Greene Publishing Associates and Wiley-Interscience (Chapter 3, In Vitro assays for Mouse Lymphocyte Function; Chapter 6, Cytokines and their cellular receptors; Chapter 7, Immunologic studies in Humans); Weinberger et al., Proc. Natl. Acad. Sci. U.S.A. 77:6091-6095, 1980; Weinberger et al., Eur. J. Immun. 11:405-411, 1981; Takai et al., J. Immunol. 137:3494-3500, 1986; Takai et al., J. Immunol. 140:508-512, 1988.

Assays for Measuring the Activity of IL-21/IL21R Agonists or Antagonists as Regulators of Hematopoiesis

IL-21/IL21R agonists or antagonists may be useful in regulation of hematopoiesis and, consequently, in the treatment of myeloid or lymphoid cell deficiencies. Even marginal biological activity in support of colony forming cells or of factor-dependent cell lines indicates involvement in regulating hematopoiesis, e.g. in supporting the growth and proliferation of erythroid progenitor cells alone or in combination with other cytokines, thereby indicating utility, for example, in treating various anemias or for use in conjunction with irradiation/chemotherapy to stimulate the production of erythroid precursors and/or erythroid cells; in supporting the growth and proliferation of myeloid cells such as granulocytes and monocytes/macrophages (i.e., traditional CSF activity) useful, for example, in conjunction with chemotherapy to prevent or treat consequent myelo-suppression; in supporting the growth and proliferation of megakaryocytes and consequently of platelets thereby allowing prevention or treatment of various platelet disorders such as thrombocytopenia, and generally for use in place of or complimentary to platelet transfusions; and/or in supporting the growth and proliferation of hematopoietic stem cells which are capable of maturing to any and all of the above-mentioned hematopoietic cells and therefore find therapeutic utility in various stem cell disorders (such as those usually treated with transplantation, including, without limitation, aplastic anemia and paroxysmal nocturnal hemoglobinuria), as well as in

repopulating the stem cell compartment post irradiation/chemotherapy, either in-vivo or ex-vivo (i.e., in conjunction with bone marrow transplantation or with peripheral progenitor cell transplantation (homologous or heterologous)) as normal cells or genetically manipulated for gene therapy.

5 The activity of the IL-21- or IL-21R binding agents may, among other means, be measured by the following methods:

Suitable assays for proliferation and differentiation of various hematopoietic lines are cited above.

10 Assays for embryonic stem cell differentiation (which will identify, among others, proteins that influence embryonic differentiation hematopoiesis) include, without limitation, those described in: Johansson et al. Cellular Biology 15:141-151, 1995; Keller et al., Molecular and Cellular Biology 13:473-486, 1993; McClanahan et al., Blood 81:2903-2915, 1993.

15 Assays for stem cell survival and differentiation (which will identify, among others, proteins that regulate lympho-hematopoiesis) include, without limitation, those described in: Methylcellulose colony forming assays, Freshney, M.G. In Culture of Hematopoietic Cells. R. I. Freshney, et al. eds. Vol pp. 265-268, Wiley-Liss, Inc., New York, N.Y. 1994; Hirayama et al., Proc. Natl. Acad. Sci. U.S.A. 89:5907-5911, 1992; Primitive hematopoietic colony forming cells with high proliferative potential, McNiece, 20 I. K. and Briddell, R. A. In Culture of Hematopoietic Cells. R. I. Freshney, et al. eds. Vol pp. 23-39, Wiley-Liss, Inc., New York, N.Y. 1994; Neben et al., Experimental Hematology 22:353-359, 1994; Cobblestone area forming cell assay, Ploemacher, R. E. In Culture of Hematopoietic Cells. R. I. Freshney, et al. eds. Vol pp. 1-21, Wiley-Liss, Inc., New York, N.Y. 1994; Long term bone marrow cultures in the presence of stromal 25 cells, Spooncer, E., Dexter, M. and Allen, T. In Culture of Hematopoietic Cells. R. I. Freshney, et al. eds. Vol pp. 163-179, Wiley-Liss, Inc., New York, N.Y. 1994; Long term culture initiating cell assay, Sutherland, H. J. In Culture of Hematopoietic Cells. R. I. Freshney, et al. eds. Vol pp. 139-162, Wiley-Liss, Inc., New York, N.Y. 1994.

The invention will be further illustrated in the following non-limiting examples.

30

Example 1: Isolation and Characterization of Murine MU-1 cDNAs:

A partial fragment of the murine homolog of the Mu-1 receptor was isolated by PCR using oligonucleotides derived from the human sequences. cDNA was prepared from RNA isolated from 17 day old murine thymus and from the murine 2D6 T cell line.

- 5 A DNA fragment of approximately 300 nucleotides was amplified from the cDNA by PCR with the following oligonucleotides, corresponding to regions 584-603 and 876-896, respectively, of the human cDNA sequence in FIG. 1 (corresponding to SEQ ID NO:1):

10 AGCATCAAG CCGGCTCCCC (5p) (SEQ ID NO:11)
CTCCATTAC TCCAGGTCCC (3p) (SEQ ID NO:12)

- Amplification was carried out using Taq polymerase in 1X Taq buffer containing 1.5 mM of magnesium chloride for 30 cycles at 94° C for one minute, 50° C for 1 minute, and 72 ° C for one minute. The DNA sequence of this fragment was determined, and two oligonucleotides were derived from an internal portion of this fragment with the
15 following sequences:

TTGAACGTGACTGRGGCCTT (5P) (SEQ ID NO:13)
TGAATGAAGTGCCTGGCTGA (3P) (SEQ ID NO:14)

- The oligonucleotides were used to amplify an internal 262 nucleotide fragment of
20 the original PCR product (corresponding to nucleotides 781-1043 of the murine cDNA sequence of figure 1, and SEQ ID NO:9) to use as a hybridization probe to screen a cDNA library isolated from the 2D6T T cell line. Filters were hybridized at 65° C using standard 5X SSC hybridization conditions and washed into SSC at 65 ° C. Twenty clones were hybridizes to the probe in a screen of 426,000 clones. DNA sequence was
25 determined from two independent clones. Full length sequence of clone #6 confirmed that it was the full-length murine homolog of human MU-1 (SEQ ID NO:9).

- The full-length nucleotide sequence of murine MU-1 is shown in figure 1 (corresponding to SEQ ID NO:9). The nucleotide sequence has a predicted leader sequence at nucleotides 407-464, coding sequence at 407-1993, and a termination codon
30 at nucleotides 1994-1997. Nucleotides 1-406 correspond to the 5' untranslated region and nucleotides 1998-2868 correspond to the 3' untranslated region.

The predicted protein sequence of murine MU-1 is shown in figure 2 (corresponding to SEQ ID NO:10). This murine MU-1 protein contains a predicted

leader sequence determined by SPScan (score= 10.1)(corresponding to amino acids 1-19 of SEQ ID NO:10), and a predicted transmembrane domain (corresponding to amino acids 237-253 of SEQ ID NO:10). Predicted signaling motifs include the following regions: Box 1: amino acids 265-274 of SEQ ID NO:10, Box 2: amino acids 310-324 of SEQ ID NO:10, six tyrosine residues at positions 281, 319, 361, 297, and 510 of SEQ ID NO:10. Potential STAT docking sites include: STAT5, EDDGYPA (SEQ ID NO:20), STAT3, YLQR.

The oligonucleotides were used to amplify an internal 262 nucleotide fragment of the original PCR product (corresponding to nucleotides 781-1043 in of the murine cDNA sequence of figure 1, and SEQ ID NO:9) to use as a hybridization probe to screen a cDNA library isolated from the 2D6 T cell line. Filters were hybridized at 65° C using standard 5 X SSC hybridization conditions and washed into SSC at 65° C. Twenty clones were isolated that hybridized to the probe in a screen of 426,000 clones. DNA sequence was determined from two independent clones. Full length sequence of clone #6 confirmed that it was the full-length murine homolog of human MU-1 (SEQ ID NO:9).

The full-length nucleotide sequence of murine MU-1 is shown in figure 1 (corresponding to SEQ ID NO:9). The nucleotide sequence has a predicted leader sequence at nucleotides 407-464, coding sequence at nucleotides 407-1993, termination codon at nucleotides 1994-1997. Nucleotides 1-406 correspond to the 5' untranslated region and nucleotides 1998-2628 correspond to the 3' untranslated region.

The predicted protein sequence of murine MU-1 is shown in figure 2 (corresponding to SEQ ID NO: 10). This murine MU-1 protein contains a predicted leader sequence determined by SPScan (score= 10.1) (corresponding to amino acids 1-19 of SEQ ID NO: 10), and a predicted transmembrane domain (corresponding to amino acids 237-253 of SEQ ID NO:10). Predicted signaling motifs include the following regions: Box 1: amino acids 265-274 of SEQ ID NO: 10, Box 2: amino acids 310-324 of SEQ ID NO: 10, six tyrosine residues at positions 281, 319, 361, 368, 397, and 510 of SEQ ID NO:10. Potential STAT docking sites include: STAT5: EDDGYPA (SEQ ID NO:20); STAT 3:YLQR.

Example 2: Comparison of Human and Murine MU-1:

The GAP algorithm was used to compare the human and murine MU-1 amino acids. A comparison of the murine and human predicted protein sequences is shown in figure 4. The amino acids were 65.267% identical using the GAP algorithm. The alignment was generated by BLOSUM62 amino acid substitution matrix (Henikoff, S. and Henikoff, J. G. (1992)). Amino acid substitution matrices from protein blocks (Proc. Natl. Acad. Sci. USA 89: 10915-10919). Gap parameters= Gap Weight: 8, Average Match= 2.9 12, Length Weight= 2, Average Mismatch= -2.003. Percent Similarity= 69.466.

A comparison of the human and murine cDNA nucleotide sequences is shown in figure 3. The DNA sequences are 66.116% identical when aligned using the GAP algorithm. Gap Parameters: Gap Weight= 50, Average Match 10.000, Length Weight= 3, Average Mismatch= 0.000. Percent Similarity= 66.198.

Both human and mouse MU-1 proteins are members of the Type 1 cytokine receptor superfamily. Evaluation of the sequence of both murine and human MU-1 reveals the presence of potential Box-i and Box-2 signaling motifs. Six tyrosine residues are present in the cytoplasmic domain, and could also be important in signaling functions of MU-1. Comparison of the sequences of MU-1 with other members of the family suggested the presence of potential docking sites for STAT 5 and STAT 3.

Example 3: Determination of STAT signaling pathways used by Human MU-1:

BAF-3 cells were engineered to express a chimeric cytokine receptor consisting of the extracellular domain of the human EPO receptor and the intracellular domain of the MU-1 receptor. BAF-3 cells that expressed huEPORJMU-1(cyto) chimeric receptors proliferated in response to human soluble EPO. These cells were analyzed to determine which STAT molecules were phosphorylated in response to EPO signaling. Briefly, control unmodified parental BAF-3 cells and EPOR/MU chimeric BAF-3 cells were rested from IL-3 containing growth medium, and restimulated with either IL-3 or EPO for 0, 15, 30 and 60 minutes. The cells were pelleted and resuspended in ice cold lysis buffer containing orthovanadate, to preserve phosphorylated tyrosines. Equal amounts of cell lysate were electrophoresed by SDS-PAGE and blotted onto nitrocellulose

membranes for western analysis. Duplicate blots were stained for phosphorylated and nonphosphorylated forms of STAT 1, 3, 5, and 6 by using antibodies specific for each form of the STAT molecule. HELA cells, non-activated and activated with alpha-interferon were used as positive controls.

- 5 These results indicated that under these specific conditions, signaling through MU-1 results in the phosphorylation of STAT 5 at all time-points tested (1=0, T=15', T=30', 1=60'). Treatment of controls or the chimeric BAF-3 cells with IL-3 resulted in phosphorylation of STAT 3, but not STAT 1 or 5.

10 **Example 4: Tissue Expression of Murine and Human MU-1**
Northern Analysis

- Northern blots of polyA+ RNA from various tissues (Clontech, Palo Alto, CA) were performed as recommended by the manufacturer. For the murine blots, a 262 nucleotide fragment corresponding to nucleotides 781-1043 of Fig. 1 and SEQ ID NO:9
15 was used for hybridization.

 A single transcript of murine MU-1 was detected in adult murine spleen, lung, and heart tissues. The larger transcript observed in human tissues was not observed in mouse tissues.

- Two transcripts of human MU-1 were detected in adult human lymphoid tissues,
20 PBLs, thymus, spleen and lymph node, and in fetal lung.

In Situ Hybridization

- In situ hybridization studies were performed by Phylogency Inc. of Columbus, OH (according to the method of Lyons et al., 1990, J. Cell. Biol: 111:2427-2436.) Briefly, serial 5-7 micron paraffin sections were deparaffinized, fixed, digested with
25 proteinase K, treated with tri-ethanolamine and dehydrated. cRNAs were prepared from linearized cDNA templates to generate antisense and sense probes. The cRNA transcripts were synthesized according to manufacturer's conditions (Ambion) and labeled with 35S-UTP. Sections were hybridized overnight, stringently washed and treated with RNAase A and dipped in nuclear track emulsion and exposed for 2-3 weeks. Control
30 sections were hybridized with sense probes to indicate the background level of the procedure. The murine probe consisted of a n 186bp fragment corresponding to

nucleotides 860-1064 (SEQ ID NO:9). The human probe was a 23 bp PCR product generated from human MU-1 DNA.

Murine MU-1 expression was observed in the lymph nodes of the adult small intestine at germinal centers and muscularis externa. Specialized lymph nodes and Peyer's patches also exhibited murine MU-1 expression.

Human MU-1 expression was detected at germinal centers of the lymph nodes in the cortex. The medulla, which contains macrophages, was negative for human MU-1. In human spleen, human MU-1 expression was detected in the regions of white pulp but not red pulp.

Example 5: Expression of Human MU-1 in cells and cell lines:

RNAse protection analysis was performed on resting and activated human T cells and the B cell lines, Raji and RPMI 8866, and the T cell line Jurkat. Human T cells were activated with anti-CD3 and anti-CD28. The cell lines were activated by Phorbol ester and ionomycin. MU-1 riboprobe-producing plasmid was constructed by inserting a 23 bp PCR product (PCR was performed by using 5' primer

CACAAAGCTTCAGTATGAGCTGCAGTACAGGAACCGGGGA (SEQ ID NO: 15) and 3' primer

CACAGGATCCCTTTAACTCCTCTGACTGGGTCTGAAAGAT (SEQ ID NO:16)) into the BamH I and HindIII sites of pGEM3zf(-) (Promega, Madison, WI) vector. To make the riboprobe, the riboprobe-producing plasmid was linearized with HindIII. The resulting DNA was phenol/chloroform extracted and precipitated with ethanol. T7 RNA polymerase was used to make the riboprobe according to the protocol suggested by the vendor (PharMingen, San Diego, CA). The RNAse protection assay was performed by using PharMingen's RiboQuant Multi-Probe Ribonuclease Protection Assay system. 2.0 µg of total RNA were included in each RPA reaction, after RNAse digestion, the protected riboprobes were run on a QuickPoint rapid nucleic acid separation system (Novex, San Diego, CA). Gels were dried and exposed according to the suggestion of the vendor.

Human MU-1 RNA is upregulated in anti-CD3 +anti-CD28 stimulated human purified CD3+ cells when compared with unstimulated populations. MU-1 is also upregulated upon restimulation in Th1 and Th2-skewed T cell populations. The B cell

lines, RPMI 8866 and Raji, constitutively express MU-1 while the Jurkat T-cell line does not.

Example 6: Binding of Human MU-1 to Known Cytokines

Both human and murine Ig fusion proteins were constructed and immobilized on Biacore chips in an effort to identify the ligand for MU-1. A variety of cell culture conditioned media as well as a panel of known cytokines were evaluated for binding to MU-1. Some cytokines were also tested in combination with other receptor chains in the family to consider the possibility that MU-1 may require a second receptor chain for ligand binding. The following cytokines were tested and found to be negative for MU-1 binding: mIL-2, hIL-2, hIL-i5, mIL-7, TSLP, TSLP+IL7, TSLP+IL7R, TSLP+IL7g, TSLP+ IL-2, TSLP + 1L2 + IL2Rbeta, IL2Rbeta, IL2Rgamma, IL7R, 1L2+2Rbeta, 1L2+2Rgamma, IL1 5+IL2Rbeta, 1L15+2Rgamma, 1L7+2Rgamma, IL2+IL7R, IL1 5+IL7R, IL7+IL7R. Known receptors have been immobilized as well and tested for MU-1 binding with negative results. IL-15 will bind to IL2Rb but not IL2Rg or MU-1.

Example 7: Effect of Modulation of IL-21/IL-21R Pathway on the Severity of Symptoms in Collagen-Induced Arthritis (CIA) Mice

This example shows that IL-21R antagonists, e.g., IL-21R-Ig fusion proteins (murine IL21RFc protein or "muIL21RFc") or anti-IL21R antibodies, ameliorate symptoms in a collagen-induced arthritis (CIA) murine model. In contrast, administration of IL-21 exacerbates the arthritic symptoms in CIA mice.

Male DBA/1 (Jackson Laboratories, Bar Harbor, Maine) mice were used for all experiments. Arthritis was induced with the use of bovine collagen type II (Chondrex, Redmond, WA). Bovine collagen type II (Chondrex, Redmond, WA) was dissolved in 0.1 M acetic acid and emulsified in an equal volume of CFA (Sigma) containing 1mg/ml *Mycobacterium tuberculosis* (strain H37RA). 100 µg of bovine collagen was injected subcutaneously in the base of the tail on day 0. On day 21, mice were injected subcutaneously, in the base of the tail, with a solution containing 100 µg of bovine collagen in 0.1M acetic acid that had been mixed with an equal volume of Incomplete Freund's adjuvant (Sigma). Naïve animals received the same sets of injections, minus collagen. The dosing protocol is shown schematically in Figure 16. MuIL21RFc was

administered prophylactically or therapeutically to DBA mice. In the therapeutic regimen, treatment was initiated if disease was observed for two consecutive days in a mouse.

Mice were monitored at least three times a week for disease progression.

- 5 Individual limbs were assigned a clinical score based on the index: 0 = normal, no swelling; 1 = visible erythema accompanied by 1-2 swollen digit, or mild swelling in ankle; 2 = pronounced erythema, characterized by mild to moderate paw swelling and/or two swollen digits; 3 = extensive swelling of the entire paw, i.e., extending into ankle or wrist joint; 4 = resolution of swelling, ankylosis of the paw; difficulty in use of limb or
10 joint rigidity. Thus, the sum of all limb scores for any given mouse yielded a maximum total body score of 16.

At various stages of disease, animals were euthanized, tissues were harvested and paws were fixed in 10% formalin for histology or 4% paraformaldehyde, pH 7.47, decalcified in 20% EDTA (pH 8.0) and embedded in paraffin for *in situ* hybridization.

- 15 Using light microscopy the paws were scored on a 5-grade scoring method (0-4) to characterize the intensity and extent of arthritis. Inflammatory infiltrates were used for scoring in addition to other changes related to the inflammation, such as pannus formation, fibrous of the synovial membrane, articular cartilage erosion and/or subchondral bone destruction. Histology grades were determined using readings of
20 individual paws: NAD=0 or nothing abnormal discovered; 1=Slight to moderate; 2: Mild to moderate; 3: Marked and 4:Massive.

- A reduction in the severity of the symptoms was observed after prophylactic treatment of CIA mice using muIL21RFc (100 µg or 200 µg) administered intraperitoneally (ip) every other day starting one day before the collagen boost (data not
25 shown).

The effects of muIL21RFc (200 µg/mouse 3x/week) on a semi-therapeutic CIA mouse as a function of day post treatment are shown in *Figure 17*. Mouse Ig (200 µg/mouse 3x/week) was used as a control. A reduction in the severity score is shown starting from day 7 post treatment.

- 30 Administration of IL-21 exacerbates the arthritic symptoms in CIA mice. *Figure 18* is a graph depicting the mean CIA score as a function of days post collagen boost.

Mice administered with 1 mg of murine IL-21 intraperitoneally (ip) showed higher CIA scores than control PBS-treated mice.

These experiments demonstrate that administration of an IL-21R antagonist, e.g., IL-21R-Fc fusion proteins, to CIA mice either prophylactically or semi-therapeutically significantly ameliorated arthritic symptoms. In contrast, administration of IL-21 exacerbates the symptoms.

Example 8: *In situ* hybridization of IL-21R transcripts

The expression of IL-21R mRNA in arthritic paws of mice with CIA was determined. Anti-sense murine IL-21R riboprobes were used (Figure 19, panel A); sense probes were used as negative controls (Figure 19, panel B).

Digoxigenin labeled probes were prepared with the use of a DIG RNA labeling mix (Roche Diagnostics, Mannheim, Germany), as described by the manufacturer. Expression of IL-21 receptor mRNA was detected in macrophages, neutrophils, fibroblasts, a subpopulation of lymphocytes, synoviocytes and epidermis (Figure 19, panel A). Decreased staining was seen in the control paws or with sense probes (Figure 19, panel B). mIL-21R mRNA positive cells were: neutrophils (N), and macrophages (M). *In situ* hybridization shows enhanced expression of IL-21R in the paws of arthritic mice.

Example 9: *In vivo* Activation of IL-21 Receptor Generates Potent Anti-Tumor Responses

This example shows that administration of IL-21-expressing tumor cells enhances anti-tumor responses *in vivo* in both immunogenic and non-immunogenic tumor models.

CD8⁺ T cells and NK cells are necessary for tumor destruction and subsequent development of tumor antigen specific T cells. IL-21 present at the tumor microenvironment potentiates both innate and adaptive immunity *in vivo* anti-tumor responses.

Experimental Procedures

Mice. Female C57BL/6, Balb/C, IFN γ ^{-/-} and IL-10^{-/-} mice in C57BL/6 background were purchased from Jackson Laboratories (Bar Harbor, ME.). C57BL/6 scid and nude mice

were purchased from Taconic (Germantown, NY). IL-21R^{-/-} mice were generated at Wyeth Research, MA. and maintained at Charles Rivers Laboratories (Andover, MA) (Kasaian, M.T. *et al.* (2002) *Immunity*. 16:559-569). Mice were maintained and treated in accordance with National Institutes of Health and American Association of Laboratory
5 Animal Care regulations.

Tumor cell lines and reagents. Two tumor cells lines, B16F1 melanoma and MethA fibrosarcoma, were used in the experiments described herein. B16F1 melanoma and MethA fibrosarcoma cells have been extensively used in tumor vaccination models to
10 study the *in vivo* anti-tumor effect of various cytokines. B16F1 tumor cells are poorly immunogenic, in that previous vaccination with irradiated wild-type tumor cells only protects 20% of vaccinated mice against subsequent live B16F1 challenge (Dranoff, G. *et al.* (1993) *PNAS* 90:3539-3543). In contrast, MethA, a methycholesterol induced fibrosarcoma, are highly immunogenic (i.e., vaccination with irradiated MethA cells
15 leads to almost 100% protection against subsequent live MethA challenge).

B16F1 melanoma cells were maintained in culture in DMEM medium supplemented with 10% heat in-activated fetal bovine serum, 2% glutamine, and 1% penicillin-streptomycin. MethA fibrosarcoma cells were maintained by intraperitoneal passage in Balb/C mice. B16F1 melanoma specific TRP-2 (SVYDFVWL, SEQ ID
20 NO: __) (van Elsas, A. *et al.* (2001) *Journal of Experimental Medicine* 194:481-489; Suttmuller, R.P. *et al.* (2001) *Journal of Experimental Medicine*. 194:823-832) and control peptide OVA₂₅₇₋₂₆₄ (SIINFEKL, SEQ ID NO: __) were synthesized at the Wyeth Research, MA. Monoclonal Abs anti-CD3, anti-CD28 and rat IgG2a, and IL-2 cytokine used in this paper were all purchased from PharMingen (San Diego, CA).

25

Generation of IL-21 and green fluorescent protein (GFP) expressing tumor cells. B16F1 and MethA tumor cells were engineered to express IL-21 and GFP, or only GFP. Retroviral vectors encoding mIL21 -IRES-GFP or IRES-GFP were constructed using GFP-RV vector (Ranganath, S. *et al.* (1998) *Journal of Immunology* 161:3822-3826).
30 High titer retrovirus was obtained by transfecting 293-VSVg ecotropic packaging cell line (Ory, D.S. *et al.* (1996) *PNAS* 93:11400-11406). Spin infections were performed at 1800 rpm for 40 minutes at room temperature. Cells were infected 3 times. Tumor cells

expressing GFP were enriched by flow sorting and the purity of GFP expressing cells was greater than 90%.

In vivo tumor studies. C57BL/6 mice were shaved on the back and injected i.d. with 10^5 B16F1-IL-21 or control B16F1-GFP. Balb/C mice were injected with either 10^6 or 2×10^6 MethA-IL-21 or control MethA-GFP cells. Tumor growth was monitored by measuring perpendicular diameters with a caliper. Mice were sacrificed when the tumors displayed severe ulceration or reached a size of 200mm^2 . In general, 10 mice per group were used in each experiment and representative results are shown from experiments repeated two or more times with similar results.

In vivo depletion studies. CD4^+ or CD8^+ T cell depletion was accomplished by i.p. injecting 400 μg per mouse of either anti-CD4 (GK1.5), anti-CD8 (53-6.7) mAbs or rat IgG isotype control for three consecutive days before tumor cell injection. Ab injections were continued every other day post tumor cell injection for 12 days. For NK cell depletion, 50 μl of rabbit anti-mouse/rat asialo GM1 polyclonal ab (Cedarlane, Ontario, CA) was injected i.p. one day before tumor cell injection. Mice similarly injected with normal rabbit serum were used for control. After tumor-cell inoculation, antibody injections were continued twice per week for two weeks. T- and NK cell-depletion was confirmed in lymph nodes and spleens one day before tumor challenge (for T cells) or on the same day of tumor cell challenge (for NK cells) by flow cytometry using relevant antibodies. FACS analysis showed that greater than 99% of the relevant population of T cells or NK cells were depleted in mice treated with anti-CD4, anti-CD8, anti asialo GM1. In contrast, mice treated with isotype controls displayed T lymphocyte profiles similar to the profiles of untreated mice.

IL-21 enzyme-linked immunosorbent assay (ELISA). Overnight supernatants from 10^6 B16F1-IL-21, B16F1-GFP, MethA-IL-21 and MethA-GFP tumor cells were assayed for IL-21 levels by ELISA as detailed in Dunussi-Joannopoulos, K. *et al.* (1998) *Blood* 91:222-230. In brief, mMu1-mutn IgG2a (Wyeth Research, MA.) was used as coating antibody and anti-mouse IL-21 (R&D systems, Minneapolis, MN.) was used as capture

Ab. Purified mIL-21 (Wyeth Research, MA) (Kasaian, M.T. *et al.* (2002) *supra*) was used as control.

5 *IL-21R mRNA expression detected by Taqman.* RNA was isolated from different tumor cell lines according to the manufacturer's instructions (Promega, Madison, WI). mRNA extracted from splenocytes that had been activated with plate bound anti-CD3 (1 μ g/ml) and anti-CD28 (1 μ g/ml) mAbs (BD PharMingen) for 24 hours was used for positive control. Purified RNA was treated with DNase (Ambion Inc, Austin, Tx) and adjusted to a concentration of 50 ng/ μ l before mRNA analysis by quantitative TaqMan polymerase
10 chain reaction (PCR) analysis. IL-21R and cyclophilin-specific primer pairs and probes were designed using PrimerExpress software and were prepared by Wyeth Research (primers: 5'-GCCTTCTCAGGACGCTATGAT-3' (SEQ ID NO:40) and 5'-CCCTACAGCACGTAGTTGGA-3' (SEQ ID NO:41) and probe TCCTGGGACTCAGCTTATGACGAACC) (SEQ ID NO:42). Standard curves for each
15 gene were generated with RNA from known IL-21R expressing cells. mRNA expression in control and transduced cell lines was normalized based on cyclophilin expression in each cell line and the results are presented as relative units (R.U.) of mRNA.

20 *Proliferation Assay.* Splenocytes (2×10^5 cells/well) from either C57BL/6 or Balb/C mice were stimulated with various concentrations of syngeneic irradiated tumor cells that expressed either GFP or IL-21 in 96-well plates. ^3H thymidine at 1 μ Ci/well (PerkinElmer Life Sciences, Boston, MA) was added during the last 6 hours of culture. After harvesting the supernatant onto glass fiber filter mats, ^3H -thymidine incorporation was determined by liquid scintillation counting.

25 *Elispot Assay.* TRP-2 specific T cell responses were determined by IFN- γ Elispot kit (R&D systems) following manufacturers instructions. In brief, 2×10^5 to 4×10^5 of splenocytes (from tumor injected mice) in 200 μ l complete medium containing 5 μ g/ml of specific TRP-2 peptides (van Elsas, A. (2001) *J. Exp. Med.* 194:481-489) or non-specific OVA peptides and 20U/ml murine IL-2 (PharMingen) were placed into each
30 well. The plate was incubated for 24 hr at 37°C in a CO₂ incubator. Plates were then incubated overnight at 4°C with detection Ab, followed by 2-hr incubation with

Streptavidin-alkaline phosphatase conjugate. Spots were visualized with 5-bromo-4-chloro-3'-indolylphosphate p-Toluidine Salt/Nitro Bluetetrazolium Chloride (BCIP/NBT) alkaline phosphatase substrate (R&D systems). Plates were washed with tap water and air-dried, and spots were counted with a stereomicroscope and re-

5 calculated to per 10^6 cells with background spots subtracted. Generally, less than 10 spots/well were detected when OVA peptide was used as antigen.

In vitro restimulation of splenocytes from tumor cell-inoculated mice. Tumor peptide specific T cell lines were generated as described in Bloom, M.B. *et al.* (1997) *J. Exp.*

10 *Med.* 185:453-459. In brief, mice were inoculated with either B16F1-GFP or B16F1-IL-21 cells. After 8-11 days, splenocytes were harvested and cultured with 5 μ g/ml of TRP-2 peptide (van Elsas, A *et al.* (2001) *supra*). On the third day of culture, 20 U/ml of IL-2 (BD PharMingen) was added to each culture. After 5 days, cells were used for ^{51}Cr release assay.

15

CTL Assay. Cytotoxicity against targets was quantified using a 4-h ^{51}Cr release assay. Briefly, RMA-S cells were pulsed with TRP-2 peptide at 10 μ g/ml and labeled with $\text{Na}_2^{51}\text{CrO}_4$ (PerkinElmer Life Sciences) for 1 hr at 37°C . After washing, ^{51}Cr labeled target cells were incubated with T cell lines generated from C57BL/6 mice injected with

20 tumor cells described earlier at different E:T ratios in 96-round bottom plates. After 4-hour incubation at 37°C , supernatants were collected and radioactivity was detected in a scintillation counter (Wallac, Turku, Finland). Percentage specific lysis was calculated as $100 \times [(\text{release by CTL} - \text{spontaneous release}) / (\text{maximal release} - \text{spontaneous release})]$. Maximal release was determined by the addition of 1% Triton X-100. Spontaneous

25 release in the absence of CTL was generally less than 15% of maximal release.

IL-21 transduced B16F1 and MethA cells secrete biologically functional IL-21

B16F1 and MethA tumor cells were transduced to express GFP plus IL-21 (B16F1/MethA-IL21) or GFP (B16F1/MethA-GFP), respectively. Overnight

30 supernatants from 10^6 of B16F1-IL-21, B16F1-GFP, MethA-IL-21 or MethA-GFP tumor cells were assayed by ELISA as described in Experimental Protocols, above. As shown in Fig. 20A, B16F1-IL-21 and MethA-IL-21 cells, but not B16F1-GFP or MethA-GFP

tumor cells, secreted substantial amount of IL-21 in overnight cultures. To determine whether the IL-21 cytokine secreted by the transduced cells was biological functional, irradiated IL-21 or GFP expressing tumor cells were used to stimulate naïve syngenic splenocytes from C56BL/6 or Balb/C mice in the presence of sub-optimal amount of anti-CD3 (500 ng/ml) and anti-CD28 (10 µg/ml). As shown in Fig. 20B and 20C, B16F1-IL-21 and MethA-IL-21 cells enhanced naïve splenocyte proliferation when compared with control GFP expressing cells at all concentrations tested. These results suggest that IL-21 secreted by transduced tumor cells is biologically functional.

10 IL-21 does not affect tumor cell growth in vitro

The effect of IL-21 on the *in vitro* growth characteristics of transduced B16F1 and MethA cells was examined. B16F1-IL-21 and B16F1-GFP or MethA-IL-21 and MethA-GFP tumor cells were cultured at a concentration of 10^5 /1.5ml culture media in 12-well plates. Cell numbers were monitored daily by trypan blue exclusion assay. The results are presented as means of duplicates. RNA from anti-CD3 and anti-CD28 mAb activated C57BL/6 splenocytes was used as positive control. Equal numbers of tumor cells were cultured in 12-well plates, and viable cell numbers were determined at various time points. As shown in Fig. 21A and 21B, the growth kinetics of IL-21 producing tumor cells was very similar to that of the GFP expressing control tumor cells in a period of 5 days. This indicates that IL-21 did not have any apparent effect on the *in vitro* growth characteristics of transduced tumor cells. The unresponsiveness of tumor cells to IL-21 was further confirmed with the lack of the IL-21R expression by tumor cells in quantitative Taqman PCR analysis. Figure 21C is a bar graph depicting quantitation of cyclophilin and IL-21R mRNA extracted from transduced tumor cells, determined by Taqman PCR. Expression of IL-21R mRNA in the transfected cells was normalized to cyclophilin values and expressed as relative unite (R.U.). RNA from anti-CD3 and anti-CD28 mAb activated C57BL/6 splenocytes was used as positive control.

30 IL-21 inhibits tumor growth in vivo

To assess the *in vivo* effects of IL-21 on tumor growth, B16F1-IL-21 or MethA-IL-21 tumor cells were inoculated i.d. into the flank of syngenic mice. As shown in Fig. 22A, no tumor formation was detected in any of the C57BL/6 mice inoculated with

B16F1-IL-21 tumor cells for more than 27 weeks post tumor injection. On the contrary, all C57BL/6 mice bearing B16F1-GFP cells grew tumors starting as early as day 9. Control tumors increased rapidly in size and these mice had to be sacrificed two to three weeks after tumor cell inoculation due to heavy tumor burden. In the MethA model, small but palpable tumor masses were detected one week post tumor inoculation with either MethA-IL-21 or MethA-GFP cells in Balb/C mice (Fig. 22B). However, MethA-IL-21 tumors gradually reduced in size starting from week two (day 11) and eventually regressed completely in 100% of mice, whereas 80% of control MethA-GFP tumors continued to grow in size until the mice were sacrificed. These results show that the presence of IL-21 at the tumor microenvironment triggers potent immune responses that lead to rejection of both immunogenic and non-immunogenic tumors.

IL-21 cannot prevent tumor growth in scid and nude mice

In order to determine the relative roles of the cellular and humoral immune responses in the rejection of IL-21-transduced tumor cells, equal numbers (10^5) of B16F1-IL-21 or control B16F1-GFP cells were injected into T and B cell deficient (scid) mice. Both B16F1-IL-21 and B16F1-GFP cells showed similar growth kinetics in scid mice (Fig. 23A), indicating that lymphocytes (B and/or T cells) are required for the IL-21 mediated tumor rejection. The role of T cells was also confirmed in experiments with C57BL/6 nude/nude mice. As shown in Fig. 23B, all nude mice inoculated with B16F1-IL-21 cells developed tumors, albeit with slower growth kinetics. Taken together, these results indicate that IL-21 mediated anti-tumor responses require the participation of the adaptive immunity.

CD8⁺ T, but not CD4⁺ T cells are required for IL-21 mediated tumor rejection

To further dissect which T cell subset(s) is (are) important for the IL-21 induced effect, *in vivo* depletion of lymphocyte subpopulations by administering anti-CD4 or anti-CD8 mAbs was performed. As shown in Fig. 25A, B16F1-IL-21 tumors did not grow in CD4⁺ T cell depleted mice and in control rat IgG treated mice. However, palpable tumors grew out in seven out of ten CD8⁺ T-cell depleted mice suggesting that CD8⁺ but not CD4⁺ T cells are necessary for the IL-21 induced anti-tumor response. Of interest, the growth of B16F1-IL-21 in CD8⁺ T cell depleted mice was significantly delayed

suggesting that cells other than CD8⁺ T-cells may be responsible for the suppression of early phase tumor growth.

NK cells are required for the IL-21 induced anti-tumor response

5 Accumulated experimental evidence supports the role of NK cells as first line of defense in promoting anti-tumor immunity (Smyth, M.J. *et al.* (2001) *International Immunology* 13:459-463; Smyth, M.J. *et al.* (2002) *Curr. Opin.Immunol.* 14:165-171; Smyth, M.J. *et al.* (2002) *Blood.* 99:1259-1266). This possibility was examined by injecting equal numbers of B16F1-IL-21 tumor cells into NK cell depleted mice or
10 control C57BL/6 mice. No detectable tumor formation was observed in control C57BL/6 mice, whereas, all of the NK cell depleted mice grew out tumors two weeks post B16F1-IL-21 tumor cell inoculation. This result shows that NK cells, in addition to CD8⁺ T cells, are necessary for IL-21 induced anti-tumor response.

15 IL-21 supports the generation of IFN γ secreting tumor antigen specific T cells and enhances tumor antigen specific CTL activity

 It has been recently demonstrated that a nonameric peptide consisting of residues 180-188 of normal melanocyte differentiation antigen tyrosinase-related protein 2 (TPR-2), is one of the tumor rejection antigens for the B16 melanoma that is recognized by B16
20 melanoma-reactive cytotoxic T lymphocytes (van Elsas, A. *et al.* (2001) *supra*; Suttmuller, R.P. (2001) *J. Exp. Med.* 194:823-832). TPR-2 peptide was used to evaluate the *in vivo* effect of IL-21 on T cell responses. Splenocytes from mice injected with either IL-21 or GFP expressing B16F1 cells were stimulated *in vitro* with TPR-2 peptide in an IFN γ ELISPOT assay. As shown in Fig. 26A, the number of IFN γ producing cells
25 in B16F1-IL-21 injected mice was 3-fold higher than that of B16F1-GFP injected mice. To further characterize IL-21 mediated anti-tumor T cell responses, splenocytes from either IL-21 or GFP expressing tumor-injected mice were first stimulated *in vitro* with TRP-2 peptide as described herein and then used for *in vitro* CTL assays. B16F1 cell
30 line used in the experiments described herein expresses low level of MHC class I molecules, as determined with flow cytometry (van Elsas, A. *et al.* (2001) *supra*; Suttmuller, R.P. (2001) *supra*; Lim, Y.S. *et al.* (1998) *Molecules & Cells* 8:629-636). Therefore, RMA-S-cells were used as antigen presenting cells in our CTL assays. As

shown in Fig. 26B, splenocytes from B16F1-IL-21 injected mice had enhanced cytolytic activity towards TRP-2 peptide pulsed RMA-S cells compared to GFP expressing tumor (Fig. 26C) at all E:T ratios, despite some cross-reactivity observed towards OVA peptide.

These results indicate that IL-21-mediated tumor rejection supports the development of tumor antigen-specific cytolytic T cell responses. TRP-2 is one of the antigens that is shared between B16F1 tumor cells and normal melanocytes, thus CTLs detected here are actually autoreactive T cells. Indeed, 10-20% of C57BL/6 mice developed local hair and skin depigmentation at the site of B16F1-IL-21 tumor cell but not control cell injection (data not shown).

10 IL-21 induced anti-tumor response is independent of IFN- γ and IL-10

IFN γ and IL-10 have been reported previously to be important for tumor rejection (Gerard, C.M. et al. (1996) *Human Gene Therapy*. 7:23-31; Lim, Y.S. et al. (1998) *Molecules & Cells* 8:629-636). ELISOP results suggest that IL-21 enhanced tumor-specific cells producing IFN γ but not IL-10. To test the involvement of IFN γ and IL-10 in IL-21-mediated anti-tumor effect, equal numbers of B16F1-IL-21 or B16F1-GFP tumor cells were injected into either IFN $\gamma^{-/-}$ or IL-10 $^{-/-}$ mice. As shown in Figs. 27A-27B, none of the aforementioned cytokine deficient mice formed tumors post B16F1-IL-21 cell injection. However, B16F1-GFP-control cells grew out tumors in all of the IFN $\gamma^{-/-}$ and IL-10 $^{-/-}$ mice. These data show that IL-21 mediated anti-tumor effect does not require active participation of IFN γ or IL-10.

Cytokine gene-therapy was used to study the *in vivo* immunoregulatory potential of IL-21. This example shows that IL-21, when secreted at the tumor site by genetically modified B16F1 melanoma or MethA fibrosarcoma cells, fuels efficacious antitumor immune responses that require the presence of the cognate IL-21R (Fig. 23). CD8 $^{+}$ T cells and NK cells are necessary for tumor destruction and subsequent development of tumor antigen specific T cells (Fig. 24A and 24B). Major Th1 and Th2 cytokines, IFN γ and IL-10 respectively, do not appear to be essential for the IL-21 mediated anti-tumor responses (Fig. 26A and 26B).

The rapid and definitive elimination of IL-21-transduced B16 melanoma tumor cells described herein is in contrast with data obtained in other cytokine gene vaccination models. Mice immunized with IL-4-, IL-5-, IL-6-, IL-12-, IFN γ -, TNF α - or GM-CSF-transduced B16 vaccines displayed moderate delay in tumor formation, albeit eventually

all mice succumbed to lethal tumors (Dranoff, G. *et al.* (1993) *PNAS* 90:3539-3543; Nagai, H. *et al.* (2000) *J. Invest. Dermat.* 115:1059-1064). Furthermore, GM-CSF, IL-5, IL-6, and TNF- α expressing cells caused significant side effects, ranging from hepatosplenomegaly, wasting and shivering, to death. To date, only IL-2 and IL-10 have
5 been reported in the literature to induce complete regression of transduced B16 tumors *in vivo* (Dranoff, G. *et al.* (1993) *supra*; Gerard, C.M. (1996) *Human Gene Therapy* 7:23-31). In the studies described herein, syngeneic mice injected with live B16F1-IL-21 or MethA-IL-21 transduced tumor cells did not develop clinically overt tumor for a period of more than 27 weeks post tumor inoculation. Furthermore, flow cytometry analysis of
10 lymphoid cells from spleens and lymph nodes removed from immunized mice did not show any major changes in cell population (data not shown), suggesting that the paracrine secretion of IL-21 at the tumor microenvironment orchestrates efficacious anti-tumor responses without causing detectable systemic side effects.

B16F1-IL-21 tumor grew out in IL-21R^{-/-} mice but not in C57BL/6 (control)
15 mice. This strongly indicates that IL-21 and IL-21R interaction is critical for the IL-21-mediated effect despite of the apparent redundancy in IL-21R signaling pathway with other cytokine receptors (Parrish-Novak *et al.* (2000) *supra*; Ozaki *et al.* (2000) *supra*; Asao *et al.* (2001) *supra*). The outgrowth of B16F1-IL-21 tumors in IL-21R^{-/-} mice, however, should not be attributed to an intrinsic defect in T and NK cell development, as
20 IL-21R^{-/-} mice have normal NK and T cells which respond to cytokines other than IL-21 (Kasaian, M.T. *et al.* (2002) *supra*).

Innate and the adaptive immune system may participate in the immune responses against tumors. IL-21 can enhance both NK and T cell activation *in vitro*. The *in vivo* depletion experiment described herein showed that both NK cells and CD8⁺ T-cells are
25 required for complete IL21-B16F1 tumor rejection. Involvement of NK cells in tumor surveillance is confirmed by the NK depletion experiment described herein (Fig. 24B). Given that B16F1-IL-21 cells express very low level of MHC class I molecules (data not shown), inoculation of live B16F1-IL-21 cells may generate an initial local inflammatory response. NK cells can act as a first line of defense against tumor cells (Trinchieri, G.
30 (1994) *J. Exp. Med.* 180:417-421; Levitsky, H.I. *et al.* (1994) *J. Exp. Med.* 179:1215-1224; Wu, T.C. (1995) *J. Exp. Med.* 182:1415-1421). This early phase of tumor killing generates a plethora of tumor derived antigens, which in turn can be taken up by

professional APCs and presented to MHC-restricted cytotoxic T cells. The contribution of NK cells to the early surveillance of B16F1-IL-21 cells before adaptive immune responses being generated is further reflected by the significantly delayed B16F1-IL-21 tumor growth in CD8⁺ T cell depleted mice (Fig. 24A).

- 5 Evidence of the involvement of T cells in IL-21 induced tumor regression is provided by the restoration of tumor growth in nude mice (Fig. 23B). Both CD4⁺ and CD8⁺ T cells have been described in various tumor cell-based vaccine strategies to be important for the induction of tumor regression and the development of protective immunity (Colombo, M.P., and G. Forni (1994) *Immunology Today* 15:48-51; Hock, H. et al. (1993) PNAS 90:2774-2778; Hung, K. et al. (1998) *J. Exp. Med.* 188:2357-2368; Segal, B.M. et al. (2002) *J. Immunol.* 168:1-4). Studies have shown that CD4⁺ T helper cells are required for the activation of naïve CD8⁺ T cells (Clarke, S.R. (2000) *Journal of Leukocyte Biology.* 67:607-614). In contrast, the findings described above demonstrate that CD8⁺ but not CD4⁺ T cells are required for the anti-tumor response of IL-21. The
- 10 findings disclosed herein appear to be consistent with recent studies that have demonstrated that eliminating CD4⁺ T cells may actually enhance the anti-tumor effect of cytokine gene therapy *in vivo* (Sutmuller, R.P. et al. (2001) *Journal of Experimental Medicine.* 194:823-832; Nagai, H. et al. (2000) *Journal of Investigative Dermatology* 115:1059-1064). IL-21 may also enhance the activation of naïve CD8⁺ T cells by
- 15 lowering the threshold of co-stimulation necessary for T cell activation, and/or augmenting antigen presentation of the IL-21R bearing APCs (Schoenberger, S.P. et al. (1998) *Nature.* 393:480-483; Bennett, S.R. et al. (1998) *Nature* 393:478-480).

- Mice injected with B16F1-IL-21 tumor cells developed hair and skin depigmentation around the injection sites. It is likely that autoimmunity against normal
- 25 mouse melanocyte was induced through a common epitope shared with B16 tumor cells in the close proximity of tumor injection sites. Using cytokine gene transduced B16 melanoma cells in conjunction with anti-CD4-depletion or CTLA-4 blockade, other investigators have also reported similar observation. However, the skin depigmentation is vitiligo-like, suggesting a more systemic involvement of the putative autoimmune cells
- 30 against melanocytes (Sutmuller, R.P. et al. (2001) *Journal of Experimental Medicine.* 194:823-832; Nagai, H. (2000) *Journal of Investigative Dermatology.* 115:1059-1064; van Elsas, A. et al. (1999) *Journal of Experimental Medicine.* 190:355-366). TRP-2 is

one of such shared epitopes between B16F1 tumor cells and normal melanocytes. As T cells with high affinity to self-antigens are deleted in the thymus, TRP-2 specific T cells escaping thymic deletion must be of low avidity. Nevertheless, B16F1-IL-21 tumor injected mice have more than three fold of TRP-2 specific T cells than B16F1-GFP (control) tumor injected mice, suggesting that IL-21 lowers the threshold for naïve T cell activation in B16F1-IL-21 injected mice.

The anti-tumor activity of IL-21 also correlates with enhanced CD8⁺ T cell functions, as demonstrated by both increased antigen specific IFN γ production, and by augmented tumor specific CTL activities. Although IFN γ and IL-10 have both been described to play an essential role in tumor rejection, our experiments with IFN γ and IL-10 deficient mice indicate that their presence is not required for IL-21 mediated tumor rejection.

In summary, the results presented herein suggest that at least three mutually nonexclusive mechanisms underlie the IL-21-mediated anti-tumor response: (1) paracrine secretion of IL-21 in the tumor microenvironment triggers and supports the initial innate tumor surveillance mediated by NK cells; (2) IL-21 activates APCs and facilitates uptake of tumor debris and subsequent tumor antigen presentation to naïve T cells, hence the adaptive immune response by CD8⁺ T cells; (3) IL-21 lowers the threshold for naïve T cell activation, thereby allowing the recruitment and activation of low-affinity autoreactive T cells that escaped central tolerance.

Example 10: IL-21 Enhances Antigen-Specific CD8⁺ T Cell Responses

This Example shows that an analysis of IL-21 and IL-21R mRNA expression, and the effects of IL-21 on the responses of, murine CD8⁺ T cells. Naïve CD8⁺ T cells express IL-21R and expression is upregulated upon stimulation. CD8⁺ T cells only express detectable IL-21 mRNA when stimulated in the presence of exogenous IL-21. IL-21 specifically enhances proliferation of antigen- or anti-CD3 mAb-stimulated CD8⁺ T cells although it is not as potent as IL-2 at limiting levels of stimulation. However, compared to IL-2, the presence of IL-21 during *in vitro* priming of CD8⁺ T cells results in the development of effector cells with increased ability to lyse target cells and produce IFN γ . These findings support a role for IL-21 in the generation of CD8⁺ T cell responses.

Experimental Protocols

Reagents. Murine IL-21 was produced by transfection of COS cells and concentration of the supernatant; activity was referenced to rmIL-21 (R&D Systems, Minneapolis, MN) by bioassay. Equivalent volumes of concentrated mock transfectant supernatant were used as a control. IL-21R.Fc was constructed linking the predicted extracellular domain of human IL-21R to murine IgG2a carboxy tail with mutations to minimize Fc binding and complement fixation. rhIL-2 (R&D Systems, Minneapolis, MN), rmIL-12 (Wyeth, Cambridge, MA), rmIL-15 (R&D Systems), rmIL-21 (R&D Systems) were used as indicated.

Mice. 2C TCR Tg mice whose TCR is specific for the Ld allo-antigen were bred and maintained at Charles River Laboratories (Wilmington, MA).

RPAs

T cell assays. T cells were purified from 2C TCR Tg lymph nodes using negative selection columns (R&D Systems) or sorted based on CD8 and CD62L expression. Purified T cells (2×10^5 /ml) were stimulated with plate-bound anti-CD3 mAb (2C11; Wyeth, Cambridge, MA) or irradiated Balb/c splenocytes (2000R; $1-5 \times 10^6$ /ml) and cytokines as indicated. Proliferation was determined by ^3H -thymidine incorporation (1 μCi /well). For priming, cultures were set as above with the indicated cytokines for 4-6 days. Cells were harvested, washed, counted and used in 4hr CTL assays with 5×10^3 ^{51}Cr -labeled P815 (specific) or EL4 (non-specific) targets at the effector:target ratios noted or restimulated on anti-CD3 mAb coated plates (1 $\mu\text{g}/\text{ml}$) for 40hrs.

IL-21R and IL-21 expression by CD8+ T cells. Previous reports have shown lymphoid-restricted expression of IL-21R and IL-21 expression by activated human CD4+ T cells. Using RNase protection assays we have analyzed the expression of IL-21R and IL-21 mRNA by naïve murine CD8+ T cells. CD8+CD62L+ 2C TCR+ cells were sorted and stimulated with anti-CD3 mAb in the presence of mock- or IL-21-transfected COS supernatant or anti-CD28 mAb and RNA was prepared for IL-21R or IL-21 RNase protection assays (Figure 27A).

Freshly isolated CD8+ T cells expressed IL-21R mRNA implying that these cells are capable of responding to IL-21. Quantitation of signal intensity relative to an internal standard showed that TCR-mediated activation resulted in ~2-fold increase in IL-21R mRNA levels (Figure 27A). TCR-activation with CD28 costimulation resulted in ~4-
5 fold increase of IL-21R mRNA, suggesting that physiologic, 2-signal activation of CD8+ T cells would result in up-regulation of IL-21R expression and increased sensitivity to IL-21. Interestingly, the presence of IL-21 during stimulation also induced a ~4-fold increase in IL-21R expression, providing evidence for a mechanism for IL-21 to amplify IL-21-responsiveness during CD8+ T cell activation. The expression of IL-21R by naïve
10 CD8+ T cells distinguishes it from receptors of other cytokines which also act on CD8+ T cells such as those for IL-2, IL-12 or IL-15 which require activation for expression. Nearly identical results were observed in non-transgenic CD8+ T cells (data not shown).

IL-21 mRNA expression was analyzed from the same purified murine CD8+ T cells. Naive, anti-CD3, or anti-CD3/anti-CD28 mAb-activated murine CD8+ T cells
15 were found not express significant IL-21 mRNA (Figure 28B). This is in agreement with results using human CD8+ T cells and contrasts with observations in CD4+ T cells where activation has been shown to up-regulate IL-21 mRNA expression. Interestingly, stimulation with anti-CD3 mAb and exogenous IL-21 resulted in strong upregulation of IL-21 mRNA expression (2-fold after 36 hours and 8-fold after 60 hours of stimulation)
20 showing that CD8+ T cells are capable of producing this cytokine.

Thus, CD8+ T cells express IL-21R mRNA, and in the presence of exogenous IL-21 up-regulate further IL-21R expression as well as de novo IL-21 expression. Activated CD4+ T cells are the likely source of the initial IL-21 driving these events. In this way, IL-21 is similar to IL-2 which is also produced predominantly by CD4+ T cells and
25 utilized by CD8+ T cells. Our results suggest a mechanism exists for maintaining IL-21 responsiveness during CD8+ T cell activation independent of CD4+ T cell-derived IL-21.

IL-21 enhances proliferation of CD8+ T cells.

IL-21 has been shown to augment the proliferation of human B cells stimulated with anti-CD40 and human and murine T cells stimulated with immobilized anti-CD3
30 mAb [1, 2]. Here, we analyzed the effect of IL-21 on antigen-stimulation of purified CD8+ murine T cells and found that proliferation of 2C CD8+ T cells was enhanced by

IL-21 in a dose-dependent manner (Figure 29A). IL-21 exhibited similar potency to IL-15, but was not as potent as IL-2 or IL-12 (Figure 29B).

Because IL-21 can affect B cell function [1], the next experiment assesses whether IL-21 was augmenting 2C CD8+ T cell proliferation by acting directly on T cells or indirectly via effects on the APCs in this system. Purified 2C CD8+ T cells were stimulated with immobilized anti-CD3 mAb in the presence of increasing amounts of IL-21 (Figure 28C). As with antigen responses, IL-21 enhanced CD8+ T cell proliferation to anti-CD3 stimulation in a dose-dependent manner. While this does not rule out a possible effect of IL-21 on APCs, it does support that this cytokine can act directly on T cells to modulate their function. Comparison of cytokine effects using anti-CD3 stimulation of CD8+ T cells revealed that IL-21 and IL-15 induce similar levels of proliferation, however IL-2 and IL-12 are more potent than IL-21 at inducing proliferation (Figure 28D).

Because the IL-21 used in the majority of these studies was supernatant concentrated from COS transfectants, an IL-21R.Fc fusion protein was used to specifically neutralize IL-21 responses. ELISAs were done to show that mIL-21R.Fc bound to mIL-21, and that it was capable of neutralizing binding of IL-21 to its receptor (data not shown). 2C CD8+ T cells were stimulated with immobilized anti-CD3 mAb and increasing amounts of IL-21 in the presence or absence of IL-21R.Fc (Figure 28C). IL-21-induced responsiveness was inhibited by the addition of soluble IL-21R.Fc showing that the enhanced proliferation observed in the presence of IL-21-transfectant supernatant was specifically due to IL-21. IL-21R.Fc had no effect on IL-2 or IL-15-mediated proliferation (data not shown).

IL-21 specifically enhanced antigen-induced proliferation of murine CD8+ T cells, on average resulting in 3-fold and 6-fold increased proliferation with antigen or anti-CD3 stimulation, respectively (n=5). This may be the result of comparatively lower levels of IL-21R expression, competition for gamma c in the case of IL-2 and IL-15, or the biological effect of this cytokine. The data shown here extends our previous results showing IL-21 enhanced proliferation of allo-antigen stimulation of unseparated T cells and underscores the potential importance of IL-21 in physiologic responses mediated by CD8+ T cells. In contrast to its ability to potentiate CD8+ T cell proliferation, IL-21 does not support the growth of murine NK cells and inhibits IL-15-mediated NK cell

growth. These results suggest that IL-21 may activate distinct signaling programs in different cell types. In agreement with results using unseparated T cells, it was found that IL-21 was most effective at augmenting the proliferation of purified CD8⁺ T cells at suboptimal levels of anti-CD3 or antigen stimulation (data not shown). This suggests that IL-21 serves as an accessory signal enhancing the sensitivity of cells responding to limiting antigen. Like IL-2, IL-21 is made by activated CD4⁺ T cells and acts on CD8⁺ T cells to enhance their proliferation and may contribute to the T-helper activity attributed to CD4⁺ T cells. Thus, although IL-21 does enhance CD8⁺ T cell proliferation it is not as potent as T-cell-derived IL-2 or macrophage-derived IL-12 suggesting non-redundant roles for these cytokines.

IL-21 enhances lytic activity of CD8⁺ effector T cells.

Having demonstrated that IL-21 enhances CD8⁺ T cell proliferation, we next examined the effects of IL-21 on CD8⁺ T cell differentiation. 2C TCR Tg CD8⁺ T cells were primed in vitro in the presence of the indicated cytokine(s) then harvested and equal numbers of cells were assayed for CTL activity (Figure 29A). The presence of IL-21 during priming resulted in the generation of CD8⁺ effector T-cells with higher lytic activity on a per cell basis than cells primed in the absence of IL-21. This increased lytic activity was even more pronounced when IL-2 was also present during priming (Figure 29B). None of these priming conditions resulted in the ability to kill non-specific target cells (data not shown). Thus, IL-21 potently enhances the development of antigen-specific CD8⁺ T cell effector function. Other cytokines such as IL-12 and IL-15 have also been shown to enhance CTL differentiation. IL-21 was compared to these cytokines and was comparable in its ability to induce potent CTLs (Figure 29B).

Previously, we have shown that IL-21 can augment IL-15-induced development of allo-antigen specific CTL and IL-15-induced NK lytic activity. The data presented here, using defined antigen-specific priming and target cells, extends this result by showing that IL-21 alone or in combination with IL-2 can enhance CTL development from naïve CD8⁺ T cells. Several mechanisms could account for this including increased sensitivity to antigen levels, increased expression of adhesion molecules or increased expression of granule enzymes. Studies are underway to examine these possibilities, preliminary data suggests that IL-21 induced higher granzyme A mRNA expression. Importantly, although IL-2 was much more potent than IL-21 in stimulating

CD8+ T cell proliferation, priming in the presence of IL-21 generated significantly more lytic activity. Despite the fact they share homology, a receptor chain and are both made by CD4+ T cells, the findings presented here underscore the unique roles of IL-2 and IL-21 and suggest an important role for IL-21 in the generation of CD8+ T cell responses.

5 IL-21 CD8+ T cell IFN γ (or IFN γ) production.

In addition to CTL activity, differentiated CD8+ effector T cells secrete high levels of IFN γ when restimulated with antigen. Therefore, we analyzed whether IL-21 present during priming modulated the ability of CD8+ T cells to produce IFN γ and compared this to other cytokines known to affect CD8+ T cell responses. 2C TCR Tg T
10 cells were primed with antigen/APCs, IL-2 and the indicated cytokine for 5 days then equal numbers of cells were restimulated with anti-CD3 mAb (Figure 29C). IL-21 treatment during CD8+ T cell priming resulted in the generation of cells producing higher titers of IFN γ than cells primed in the absence of IL-21 (13-fold increase). Using cytokine doses that generated equivalent CTL activity, CD8+ T cells primed in the
15 presence of IL-12 or IL-21 produced similar titers of IFN γ . IL-15 was most effective at inducing IFN γ producing CD8+ T cells. As both IL-15 and IL-21 similarly enhanced the development of CD8+ T cell lytic activity (Figure 29B), this differential ability to induce IFN γ production suggests non-redundant roles for these related cytokines. These results are particularly interesting as IL-21 has recently been shown to antagonize IL-12-
20 induced IFN γ production by CD4+ T cells. Thus, IL-21 plays opposing roles in the differentiation of CD4+ and CD8+ T cells into IFN γ -producing cells. This is consistent with the ability of CD8+ T cells to produce IFN γ in an IL-12/Stat4 independent manner. In conclusion, exposure to IL-21 results in increased ability to secrete IFN γ consistent with its role as differentiation factor for CD8+ T cells.

25 The experiments described herein show that IL-21 is a potent growth and differentiation factor for CD8+ T cells responding to antigen. IL-21R-deficient animals have functional CD8+ T cells. This finding, coupled with the data presented herein, implies that IL-21 responses are not required for CD8+ T cell development/maintenance but rather that IL-21 augments the generation of effector function. Like IL-2, IL-21 is
30 made by activated CD4+ T cells and our results comparing the actions of these two cytokines suggest that although IL-2 potently drives the proliferation of CD8+ T cells, IL-21 is much more effective at promoting CD8+ T effector differentiation. Recently,

IL-21 has been shown to be produced primarily by Th2 cells, suggesting a mechanism for the development IFN γ -producing CTLs during Th2 responses. IL-15 is similar to IL-21 in its ability to induce naïve cell proliferation and CTL development but is much more effective at eliciting IFN γ production. Like IL-2, IL-12 also induces strong CD8+ T cell proliferation and like IL-21 it induces potent CTL activity and IFN γ production. Thus IL-2, IL-12, IL-15 and IL-21 are effective at enhancing different aspects of naïve CD8+ T cell responses. In addition, these cytokines are made by different cell types: IL-2 and IL-21 are secreted by activated CD4+ T cells while IL-12 is made by activated macrophages and IL-15 is produced by stromal cells, endothelial cells as well as macrophages. Taken together, this suggests that IL-2, IL-12, IL-15 and IL-21 act at distinct phases and sites of immune responses are not redundant. We have previously proposed that IL-21 may be a mediator of the transition between innate and adaptive immune responses, the data presented here showing enhancement of CD8+ T cell responses support this hypothesis. Given its potent effects on CD8+ T cells, it will be interesting to analyze the role of IL-21 in CD8+ responses to tumors and viral infections *in vivo*.

Example 11: Primary Macrophages Express IL-21R and Respond to IL-21 by Proliferating and Secreting Increased Levels of Cytokines and Chemokines

This Example describes the finding that IL-21 acts as a proliferation factor for hematopoietic cells specifically promoting the growth of macrophages and bone marrow progenitor cells.

The effect of IL-21 alone and in combination with other factors in modulating hematopoietic cell growth was examined. It was found that IL-21 acts as a proliferation factor for mouse and human hematopoietic cells specifically promoting the growth of macrophage colonies in methylcellulose from lineage negative murine bone marrow progenitors. IL-21 also acts to enhance proliferation of human CD34+ cells and results in increased expression of the macrophage specific marker CD14. Utilizing CD14+ cells sorted from human bone marrow, it has been shown by RNase Protection Analysis (RPA) that the IL-21R is expressed on macrophages, and is upregulated by TNF and IL-1 following growth in GM-CSF. Thus, suggesting a role of other inflammatory cytokines in regulating IL-21 responsiveness. Expression of IL-21R protein on mouse and human macrophages has been detected using FACS analysis. IL-21 acts directly on these cells

resulting in activation, growth and increased secretion of chemokines and cytokines. Taken together, these results show that the IL-21/IL-21R complex is important in regulating macrophage growth and function, and in influencing macrophage responses at inflammatory sites.

- 5 The contents of all references, pending patent applications and published patents, cited throughout this application are hereby expressly incorporated by reference.

Equivalents

- 10 Those skilled in the art will recognize, or be able to ascertain using no more than routine experimentation, many equivalents of the specific embodiments of the invention described herein. Such equivalents are intended to be encompassed by the following claims.

What is claimed is:

1. A method of treating or preventing an arthritic disorder, in a subject, comprising, administering to the subject an IL-21/IL-21R antagonist, in an amount sufficient to
5 inhibit or reduce immune cell activity in the subject, thereby treating or preventing the arthritic disorder.
2. A method of treating or preventing an arthritic disorder, in a subject, comprising, administering to the subject an IL-21/IL-21R antagonist in combination with another
10 therapeutic agent, in an amount sufficient to inhibit or reduce immune cell activity in the subject, thereby treating or preventing the arthritic disorder.
3. The method of claim 2, wherein the therapeutic agent is selected from the group consisting of a cytokine inhibitor, a growth factor inhibitor, an immunosuppressant, an
15 anti-inflammatory agent, a metabolic inhibitor, an enzyme inhibitor, a cytotoxic agent, and a cytostatic agent.
4. The method of claim 2, wherein the therapeutic agent is selected from the group consisting of a TNF antagonist, an IL-12 antagonist, an IL-15 antagonist, an IL-17
20 antagonist, an IL-18 antagonist, an IL-22 antagonist, a T cell depleting agent, a B cell depleting agent, methotrexate, leflunomide, sirolimus (rapamycin) or an analog thereof, a Cox-2 inhibitor, a cPLA2 inhibitor, an NSAID, and a p38 inhibitor.
5. The method of claim 4, wherein the TNF antagonist is a soluble fragments of a TNF
25 receptor.
6. The method of claim 4, wherein the TNF antagonist is a p75 human TNF receptor-IgG fusion protein.
- 30 7. The method of claim 4, wherein the therapeutic agent is an IL-15 antagonist.
8. The method of claim 4, wherein the therapeutic agent is methotrexate or leflunomide.

9. The method of claim 4, wherein the rapamycin analog is CCI-779.

10. The method of any of claims 1-9, wherein the arthritic disorder is selected from the
5 group consisting of rheumatoid arthritis, juvenile rheumatoid arthritis, osteoarthritis, psoriatic arthritis, and ankylosing spondylitis.

11. The method of claim 10, wherein the arthritic disorder is rheumatoid arthritis.

10 12. The method of claim 10, wherein the arthritic disorder is selected from the group consisting of juvenile rheumatoid arthritis, osteoarthritis, psoriatic arthritis, and ankylosing spondylitis.

13. The method of either of claim 1 or 2, wherein the subject is a mammal.

15

14. The method of claim 13, wherein the subject is a human.

15. The method of any of claim 1-14, wherein the IL-21/IL-21R antagonist is an anti-IL21R or anti-IL21 antibody, or an antigen-binding fragment thereof.

20

16. The method of any of claim 1-14, wherein the IL-21/IL-21R antagonist comprises a fragment of a human IL-21 receptor (IL-21R) polypeptide.

17. The method of claim 16, wherein the fragment comprises the extracellular domain of
25 human IL-21R and an Fc fragment of a human immunoglobulin (Ig).

18. The method of claim 17, wherein the extracellular domain of human IL-21R comprises about amino acids 20-235 of SEQ ID NO:2.

30 19. The method of claim 18, wherein the extracellular domain of human IL-21R comprises about amino acids 1-235 of SEQ ID NO:2.

20. The method of claim 19, wherein the extracellular domain of human IL-21R comprises about amino acids 1-235 of SEQ ID NO:2 and an Fc fragment of a human IgG1.
- 5 21. The method of claim 17, wherein the Fc fragment does not bind to an Fc receptor.
22. The method of claim 16, wherein the IL-21/IL-21R antagonist comprises the amino acid sequence of SEQ ID NO:29 (Figures 10A-10C), or a sequence at least 90% or more identical thereto.
- 10 23. The method of claim 16, wherein the IL-21/IL-21R antagonist comprises the amino acid sequence of SEQ ID NO:25 (Figures 8A-8C), or a sequence at least 90% or more identical thereto.
- 15 24. A method of treating or preventing a cancer or an infectious disorder, in a subject, comprising, administering to the subject an IL-21/IL-21R agonist, in an amount sufficient to increase immune cell activity, thereby treating or preventing said disorder.
- 20 25. The method of claim 24, wherein the immune cell is a CD8+ cell, and the activity is increased effector cell activity.
26. The method of claim 24, wherein the cancer is selected from the group consisting of a solid tumor, a soft tissue tumor, and a metastatic lesion.
- 25 26. The method of claim 24, wherein the infectious disorder is a bacterial, viral or parasitic infection.
27. The method of claim 24, wherein the IL-21/IL-21R agonist is administered alone.
- 30 28. The method of claim 24, wherein the IL-21/IL-21R agonist is administered in combination with an antigen from a cancer or tumor cell, or a pathogen.

29. A method for increasing the ability of a vaccine composition containing an antigen to elicit a protective immune response in a subject against the antigen, comprising administering to the subject, either simultaneously with or sequentially, to the vaccine composition, an effective adjuvanting amount of an IL-21/IL-21R agonist, such that the ability of the vaccine composition to elicit the protective immune is increased.
30. The method of claim 29, wherein the antigen is from a pathogen selected from the group consisting of a virus, bacterium, and protozoan.
31. The method of claim 29, wherein the antigen is from a cancer or tumor cell antigen.
32. The method of claim 31, wherein the antigen is expressed on the surface of a cancer cell.
33. The method of any of claim 24-32, wherein the IL-21/IL-21R agonist is administered in combination with one or more cytokines chosen from IL-2, GM-CSF or IL-15.
34. The method of any of claim 24-32, wherein the subject is a mammal.
35. The method of claim 34, wherein the subject is a human.
36. The method of any of claim 24-35, wherein the IL-21/IL-21R agonist is a human IL-21 polypeptide comprising the amino acid sequence shown as SEQ ID NO:19, or a sequence at least 90% or more identical thereto.
37. A method for modulating the activity of an immune cell chosen from one or more of a mature CD8+ T cell, CD4+ T cell, mature NK cell, macrophage or megakaryocyte, comprising contacting the immune cell, with an IL-21/IL-21R agonist or antagonist, in an amount sufficient to modulate immune cell activity.
38. A fusion protein comprising an extracellular domain of human IL-21R and an Fc fragment of a human immunoglobulin (Ig).

39. The fusion protein of claim 38, wherein the extracellular domain of human IL-21R comprises about amino acids 20-235 of SEQ ID NO:2.
- 5 40. The fusion protein of claim 38, wherein the extracellular domain of human IL-21R comprises about amino acids 1-235 of SEQ ID NO:2.
41. The fusion protein of claim 38, wherein the extracellular domain of human IL-21R comprises about amino acids 1-235 of SEQ ID NO:2 and an Fc fragment of a human
10 IgG1.
42. The fusion protein of claim 38, wherein the Fc fragment does not bind to an Fc receptor.
- 15 43. The fusion protein of claim 38, wherein the IL-21/IL-21R antagonist comprises the amino acid sequence of SEQ ID NO:29 (Figures 10A-10C), or a sequence at least 90% or more identical thereto.
44. The fusion protein of claim 38, wherein the IL-21/IL-21R antagonist comprises the
20 amino acid sequence of SEQ ID NO:25 (Figures 8A-8C), or a sequence at least 90% or more identical thereto.
45. A host cell comprising a nucleotide sequence encoding the fusion protein of claim 38.
- 25 46. An vector comprising a nucleotide sequence encoding the fusion protein of claim 38.
47. A process for producing the fusion protein of claim 38, comprising, (a) growing a culture of the host cell of the present invention in a suitable culture medium; and (b) purifying the fusion protein from the culture.
- 30 48. A pharmaceutical composition comprising an IL-21/IL-21R antagonist and a therapeutic agent selected from the group consisting of a cytokine inhibitor, a growth

factor inhibitor, an immunosuppressant, an anti-inflammatory agent, a metabolic inhibitor, an enzyme inhibitor, a cytotoxic agent, and a cytostatic agent, in a pharmaceutically acceptable carrier.

- 5 49. The pharmaceutical composition of claim 48, wherein the therapeutic agent is selected from the group consisting of a TNF antagonist, an IL-12 antagonist, an IL-15 antagonist, an IL-17 antagonist, an IL-18 antagonist, an IL-22 antagonist, a T cell depleting agent, a B cell depleting agent, methotrexate, leflunomide, sirolimus (rapamycin) or an analog thereof, a Cox-2 inhibitor, a cPLA2 inhibitor, an NSAID, and a
- 10 p38 inhibitor.
50. The pharmaceutical composition of claim 49, wherein the TNF antagonist is a soluble fragments of a TNF receptor.
- 15 51. The pharmaceutical composition of claim 50, wherein the TNF antagonist is a p75 human TNF receptor-IgG fusion protein.
52. The pharmaceutical composition of claim 50, wherein the therapeutic agent is an IL-15 antagonist.
- 20 53. The pharmaceutical composition of claim 50, wherein the therapeutic agent is methotrexate or leflunomide.
54. The pharmaceutical composition of claim 50, wherein the rapamycin analog is CCI-
- 25 779.
55. The pharmaceutical composition of any of claim 50-54, wherein the IL-21/IL-21R antagonist is an anti-IL-21- or an anti-IL-21R antibody, or a fragment of a human IL-21R.

56. The pharmaceutical composition of claim 55, wherein the IL-21/IL-21R antagonist comprises the extracellular domain of human IL-21R and an Fc fragment of a human immunoglobulin (Ig).

5 57. The pharmaceutical composition of claim 56, wherein the extracellular domain of human IL-21R comprises about amino acids 20-235 of SEQ ID NO:2.

58. The pharmaceutical composition of claim 56, wherein the extracellular domain of human IL-21R comprises about amino acids 1-235 of SEQ ID NO:2.

10

59. The pharmaceutical composition of claim 56, wherein the extracellular domain of human IL-21R comprises about amino acids 1-235 of SEQ ID NO:2 and an Fc fragment of a human IgG1.

15 60. The pharmaceutical composition of claim 59, wherein the Fc fragment does not bind to an Fc receptor.

61. The pharmaceutical composition of claim 50, wherein the IL-21/IL-21R antagonist comprises the amino acid sequence of SEQ ID NO:29 (Figures 10A-10C), or a sequence
20 at least 90% or more identical thereto.

61. The pharmaceutical composition of claim 50, wherein the IL-21/IL-21R antagonist comprises the amino acid sequence of SEQ ID NO:25 (Figures 8A-8C), or a sequence at
25 least 90% or more identical thereto.

25

62. A pharmaceutical composition useful as a vaccine, comprising an antigen from a pathogenic microorganism selected from the group consisting of a viral, bacterial and parasitic microorganism, and an effective adjuvanting amount of an IL-21/IL-21R agonist, in a pharmaceutically acceptable carrier.

30

63. A pharmaceutical composition comprising a cancer cell- or tumor cell-antigen in combination with an IL-21/IL-21R agonist, in a pharmaceutically acceptable carrier.

FIG.1

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1  GTCGACGCGG-EGGTACCAAC TGTCTGCCCA CTTCTCTCTGT GGTGTGCCTC
51  ACGGTCACTT GCTTGTCTGA CCGCAAGTCT GCCCATCCCT GGGGCAGCCA
101 ACTGGCCTCA GCCCGTGCCC CAGGCGTGCC CTGTCTCTGT CTGGCTGCCC
151 CAGCCCTACT GTCTTCTCTT GTGTAGGCTC TGCCCAGATG CCCGCTGGT
201 CCTCAGCCTC AGGACTATCT CAGCAGTGAC TCCCCTGATT CTGGACTTGC
251 ACCTGACTGA ACTCTGCCC ACCTCAAACC TTCACCTCCC ACCACCACCA
301 CTCCGAGTCC CGCTGTGACT CCCACGCCCA GGAGACCACC CAAGTGCCCC
351 AGCCTAAAGA ATGGCTTTCT GAGAAAGACC CTGAAGGAGT AGGTCTGGGA
401 CACAGCATGC CCCGGGGCCC AGTGGCTGCC TTA CTCTCTGC TGATTCTCCA
451 TGGAGCTTGG AGCTGCCTGG ACCTCACTTG CTACACTGAC TACCTCTGGA
501 CCATCACCTG TGTCTTGAG ACACGGAGCC CCAACCCAG CATACTCAGT
551 CTCACCTGGC AAGATGAATA TGAGGAACCT CAGGACCAAG AGACCTTCTG
601 CAGCCTACAC AGGTCTGGCC ACAACACCAC ACATATATGG TACACGTGCC
651 ATATGCGCTT GTCTCAATTC CTGTCCGATG AAGTTTTCAT TGTCAATGTG
701 ACGGACCAGT CTGGCAACAA CTCCCAAGAG TGTGGCAGCT TTGTCTTGGC
751 TGAGAGCATC AAACCAGCTC CCCCCTTGAA CGTGACTGTG GCCTTCTCAG
801 GACGCTATGA TATCTCTTGG GACTCAGCTT ATGACGAACC CTCCA ACTAC
851 GTGCTGAGGG GCAAGCTACA ATATGAGCTG CAGTATCGGA ACCTCAGAGA
901 CCCCTATGCT GTGAGGCCGG TGACCAAGCT GATCTCAGTG GACTCAAGAA
951 ACGTCTCTCT TCTCCCTGAA GAGTTCCACA AAGATTCTAG CTACCAGCTG
1001 CAGGTGCGGG CAGCGCCDCA GCCAGGCACT TCATT CAGGG GGACCTGGAG
1051 TGAGTGGAGT GACCCCGTCA TCTTTCAGAC CCAGGCTGGG GAGCCCGAGG
1101 CAGGCTGGGA CCCTCACATG CTGCTGCTCC TGGCTGTCTT GATCATTTGC
1151 CTGGTTTTCA TGGGTCTGAA GATCCACCTG CCTTGGAGGC TATGGAAAAA
1201 GATATGGGCA CCAGTGCCCA CCCCTGAGAG TTTCTTCCAG CCCCTGTACA
1251 GGGAGCACAG CGGGAACCTC AAGAAATGGG TTAATACCCC TTTCACGGCC
1301 TCCAGCATAG AGTTGGTGCC ACAGAGTTCC ACAACAACAT CAGCCTTACA
1351 TCTGTCACTG TATCCAGCCA AGGAGAAGAA GTTCCCGGGG CTGECGGGTC
1401 TGGAAAGAGCA ACTGGAGTGT GATGGAATGT CTGAGCCTGG TCACTGGTGC

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FIG.1 (continued)

1451 ATAATCCCCT TGGCAGCTGG CCAAGCGGTC TCAGCCTACA GTGAGGAGAG
1501 AGACCGGCCA TATGGTCTGG TGTCCATTGA CACAGTGA CT GTGGGAGATG
1551 CAGAGGGCCT GTGTGTCTGG CCCTGTAGCT GTGAGGATGA TGGCTATCCA
1601 GCCATGAACC TGGATGCTGG CCGAGAGTCT GGCCCTAATT CAGAGGATCT
1651 GCTCTTGGTC ACAGACCCTG CTTTTCTGTC TTGCGGCTGT GTCTCAGGTA
1701 GTGGTCTCAG GCTTGGAGGC TCCCCAGGCA GCCTACTGGA CAGGTTGAGG
1751 CTGTCATTTG CAAAGGAAGG GGACTGGACA GCAGACCCAA CCTGGAGAAC
1801 TGGGTCCCCA GGAGGGGGCT CTGAGAGTGA AGCAGGTTCC CCCCCTGGTC
1851 TGGACATGGA CACATTTGAC AGTGGCTTTG CAGGTTGAGA CTGTGGCAGC
1901 CCCGTGGAGA CTGATGAAGG ACCCCCTCGA AGCTATCTCC GCCAGTGGGT
1951 GGTGAGGACC CCTCCACCTG TGGACAGTGG AGCCCAGAGC AGCTAGCATA
2001 TAATAACCAG CTATAGTGAG AAGAGGCCTC TGAGCCTGGC ATTTACAGTG
2051 TGAACATGTA GGGGTGTGTG TGTGTGTGTG TGTGTGTGTG TGTGTGTGTG
2101 TGTGTGTGTG TGTGTGTGTG TGTCTTGGGT TGTGTGTTAG CACATCCATG
2151 TTGGGATTTG GTCTGTTGCT ATGTATTGTA ATGCTAAATT CTCTACCCAA
2201 AGTTCTAGGC CTACGAGTGA ATTCTCATGT TTACAAACTT GCTGTGTAAA
2251 CCTTGTTCTT TAATTTAATA CCATTGGTTA AATAAAATTG GCTGCAACCA
2301 ATTACTGGAG GGATTAGAGG TAGGGGGCTT TTGAGTTACC TGTTTGGAGA
2351 TGGAGAAGGA GAGAGGAGAG ACCAAGAGGA GAAGGAGGAA GGAGAGGAGA
2401 GGAGAGGAGA GGAGAGGAGA GGAGAGGAGA GGAGAGGAGA GGAGAGGAGA
2451 GGCTGCCGTG AGGGGAGAGG GACCATGAGC CTGTGGCCAG GAGAAACAGC
2501 AAGTATCTGG GGTACACTGG TGAGGAGGTG GCCAGGCCAG CAGTTAGAAG
2551 AGTAGATTAG GGGTGACCTC CAGTATTTGT CAAAGCCAAT TAAAATAACA
2601 AAAAAAAAAA AAAAGCGGCC GCTCTAGA

FIG.2A

1 MPRGPVAALL LLILHCAWSC LDLCYTDYL WTITCVLETR SPNPSILSLT
51 WQDEYEELQD QETFCSLHRS GHNTTHIWYT CHMRLSQFLS DEVFIVNVT
101 QSGNNSQECG SFVLAESIKP APPLNVTVAF SGRYDISWDS AYDEPSNYVL
151 RGKLQYELQY RNL RDPYAVR PVTKLISVDS RNVSLLP EEF HKDSSYQLQV
201 RAAPQPGTSF RGT WSEWS DP VIFQTQAGEP EAGWDPHMLL LLAVLIIVLV
251 FMGLKIHLPW RLWKKIWA PV PTPESFFQPL YREHSGNFKK WVNTPFTASS
301 IELVPQSSTT TSALHLSLYP AKEKKFPGLP GLEEQL ECDG MSEPGHWCII
351 PLAAGQAVSA YSEERDRPYG LVSIDTVTVG DAEGLCVWPC SCEDDGYPAM
401 NLDAGRESGP NSEDLLLVT D PAF LSCGCVS GSGLRLGGSP GSLLDRLRLS
451 FAKEGDWTAD PTWRTGSPGG GSESEAGSPP GLDMDTFD SG FAGSDCGSPV
501 ETDEGP PRSY LRQWVVRTPP PVDSGAQSS

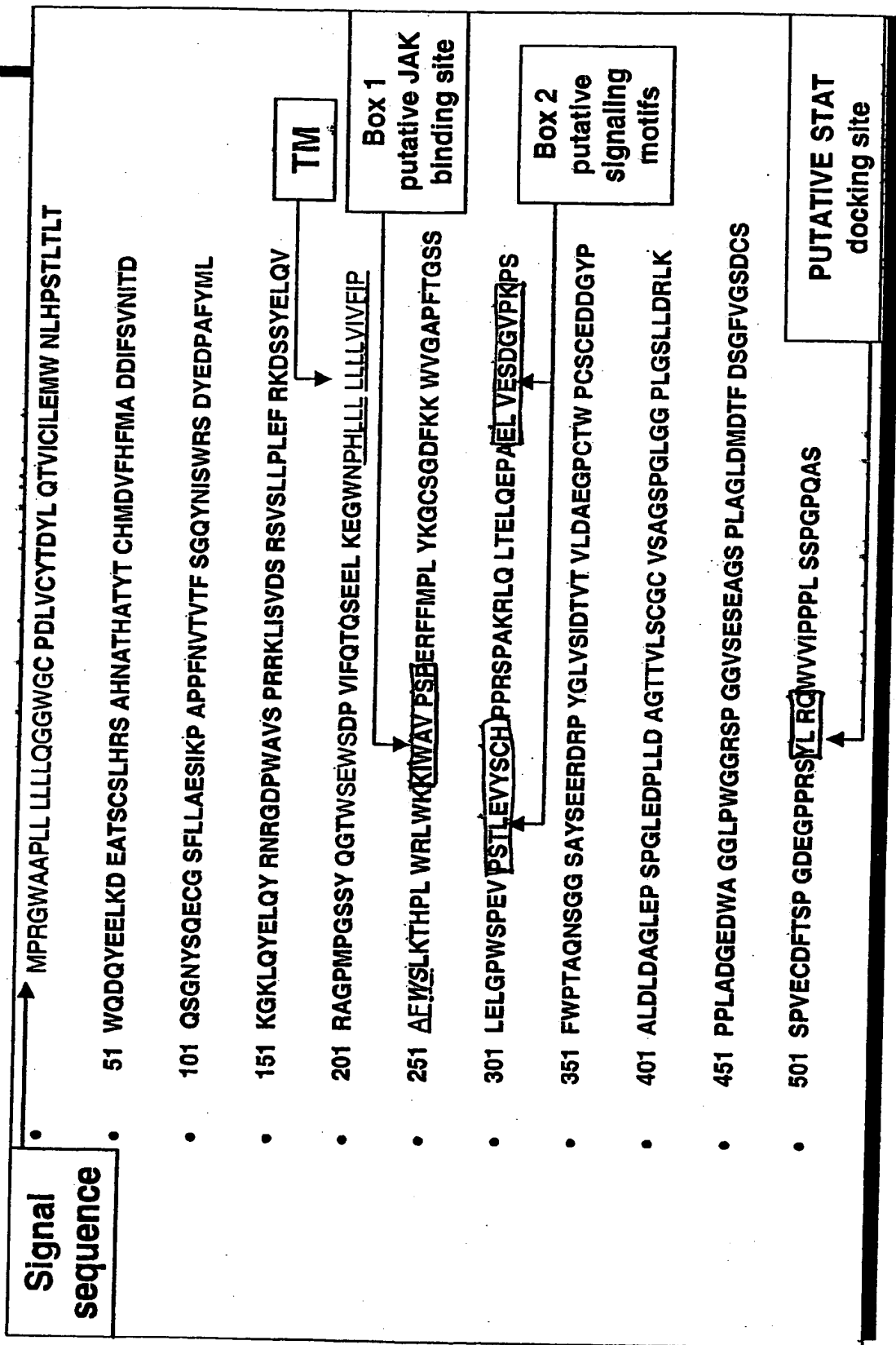


Figure 2B

FIG.3

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huMU-1 .....NNGTCGACTGGAGGCCAGCTGCCGGTCATCA 32
           ::|| | ||||| ||||| |
murMU-1 CAGCCCTACTGTCTTCCTCTGTGTAGGCTCTGCCAGATGCCCGGC.... 196

huMU-1 GAGTGACAGGTCTTATGACAGCCTGATTGGTGACTCGGGCTGGGTGTGGA 82
           || | || | || | || | ||||| || | |||||
murMU-1 TGGTCCTCAGCCTCAGGACTATCTCAGCAGTGACTC.CCCTGATTCTGGA 245

huMU-1 TTCTCACCCAGGCCCTCTGCCTGCTTTCTCAGACCCCTCATCT...GTCAC 129
           | |||| | || | | ||||| ||||| |||||
murMU-1 CTTGCACCTGACTGAACTCCTGCCACCTCAAACCTTCACCTCCCACCAC 295

huMU-1 CCCCACGCTGAACCCAGCTG.....CCACCCCAAGAAGCCCATCAGACT 173
           | |||| | || | ||||| ||||| ||||| ||||| | |
murMU-1 CACCACTCCGAGTCCCGCTGTGACTCCCACGCCAGGAGACCACCCAGT 345

huMU-1 GCCCCAGCACACGGAATGGATTCTGAGAAAGAAGCCGAAACAGAAGGC 223
           | ||||| | ||||| ||||| ||||| | || | || |||
murMU-1 G.CCCCAGCCTAAAGAATGGCTTTCTGAGAAAGACCCTGAAGGAGTAGGT 394

huMU-1 CCGTGGGAGTCAGCATGCCCGGTGGCTGGGCGGCCCTTGCTCCTGCTG 273
           | |||| | ||||| || || | || | || | ||||| |||
murMU-1 C..TGGGACACAGCATGCCCGGGGCCAGTGCTGCCTTACTCCTGCTG 442

huMU-1 CTGCTCCAGGAGGCTGGGGCTGCCCGACCTCGTCTGCTACACCGATTA 323
           | |||| |||| | || ||||| ||||| ||||| || ||
murMU-1 ATTCTCCATGGAGCTTGGAGCTGCCTGGACCTCACTTGCTACACTGACTA 492

huMU-1 CCTCCAGACGGTCATCTGCATCCTGGAAATGTGGAACCTCCACCCAGCA 373
           |||| || | || | || ||||| | || | || | ||||| |||
murMU-1 CCTCTGGACCATCACCTGTGTCTTGGAGACACGGAGCCCCAACCCAGCA 542

huMU-1 CGCTCACCCCTTACCTGGCAAGACAGTATGAAGAGCTGAAGGACGAGGCC 423
           |||| || ||||| ||||| | |||| || || ||||| |||
murMU-1 TACTCAGTCTACCTGGCAAGATGAATATGAGGAACCTCAGGACCAAGAG 592

huMU-1 ACCTCCTGCAGCCTCCACAGGTGGGCCACAAATGCCACGATGCCACCTA 473
           |||| ||||| ||||| || ||||| ||||| ||||| ||
murMU-1 ACCTTCTGCAGCCTACACAGGTCTGGCCACAACACCACACATATATGGTA 642

huMU-1 CACCTGCCACATGGATGTATTCCACTTCATGGCCGACGACATTTTCAGTG 523
           |||| |||| || | | || || || || ||||| || ||||| ||
murMU-1 CACGTGCCATATGCGCTGTCTCAATTCTGTCCGATGAAGTTTTTCATTG 692

huMU-1 TCAACATCACAGACAGTCTGGCAACTACTCCAGGAGTGTGGCAGCTTT 573
           |||| | || ||||| ||||| ||||| ||||| ||||| |||||
murMU-1 TCAATGTGACGGACAGTCTGGCAACAACTCCCAAGAGTGTGGCAGCTTT 742

huMU-1 CTCCTGGCTGAGAGCATCAAGCCGGCTCCCCCTTTCAACGTGACTGTGAC 623
           ||||| ||||| || ||||| || ||||| || ||||| |||||
murMU-1 GTCCTGGCTGAGAGCATCAACCCAGCTCCCCCTTGAACTGACTGTGGC 792

huMU-1 CTTCTCAGGACAGTATAATATCTCCTGGCGCTCAGATTACGAAGACCTG 673
           ||||| ||||| || ||||| ||||| ||||| ||||| |||||
murMU-1 CTTCTCAGGACGCTATGATATCTCCTGGGACTCAGCTTATGACGAACCT 842

huMU-1 CCTTCTACATGCTGAAGGGCAAGCTTCAGTATGAGCTGCAGTACAGGAAC 723
           || |||| ||||| ||||| || ||||| ||||| ||||| |||||
murMU-1 CCAACTAGGTGCTGAGGGGCAAGCTACAATATGAGCTGCAGTATCGGAAC 892

huMU-1 CGGGGAGACCCCTGGGCTGTGAGTCCGAGGAGAAAGCTGATCTCAGTGGA 773
           | ||||| ||||| ||||| || || ||||| ||||| |||||
murMU-1 CTCAGAGACCCCTATGCTGTGAGGCCGGTGACCAAGCTGATCTCAGTGGA 942

huMU-1 .CTCAAGAAGTGTCTCCCTCCTCCCCCTGGAGTTCCGCAAGACTCGAGCT 823

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FIG.3 (continued)

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murMU-1  ||||| ||||| || ||||| ||||| ||||| || |||||
CTCAAGAAACGTCTCTCTTCTCCCTGAAGAGTTCCACAAAGATTCTAGCT 992

huMU-1  ATGAGCTGCAGGTGCGGGCAGGGCCCATGCCTGGCTCCTCTACCAGGGG 873
| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

murMU-1  ACCAGCTGCAGGTGCGGGCAGCGCCTCAGCCAGGCATTCTATTAGGGGG 1042

huMU-1  ACCTGGAGTGAATGGAGTGACCCGGTCATCTTTAGACCCAGTCAGAGGA 923
||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

murMU-1  ACCTGGAGTGAATGGAGTGACCCGGTCATCTTTAGACCCAGGCTGGGGA 1092

huMU-1  GTTAAAGGAAGGCTGGAACCTCACCTGCTGCTTCTCCTCCTGCTTGTC 973
||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

murMU-1  GCCCAGGAGGCTGGGACCTCACATGCTG...CTGCTCCTGGCTGTCT 1139

huMU-1  TAGTCTTCATTCTGCCTTCTGGAGCCTGAAGACCCATCCATTGTGGAGG 1023
| ||| | ||||| ||||| ||||| ||||| ||||| |||||

murMU-1  TGATCATTGTCTGCTTCTCATGGGTCTGAAGATCCACCTGCCTTGGAGG 1189

huMU-1  CTATGGAAGAAGATATGGG...CCGTCCCCAGCCCTGAGCGGTTCTTCAT 1070
||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

murMU-1  CTATGGAAGAAAGATATGGGCACCACTGCCACCCCTGAGAGTTTCTTCCA 1239

huMU-1  GCCCTGTACAAGGGCTGCAGCGGAGACTTCAAGAAATGGGTGGGTGCAC 1120
||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

murMU-1  GCCCTGTACAGGAGCACAGCGGGAACCTTCAAGAAATGGGTGAATACCC 1289

huMU-1  CCTTCACTGGCTCCAGCCTGGAGCTGGGACCCCTGGAGCCAGAGGTGCC 1170
| ||||| | ||||| | ||||| ||||| ||||| ||||| |||||

murMU-1  CTTTACGGCCTCCAGCATAGAGTTGGTGCCACAGAGTTCCACAACAACA 1339

huMU-1  TCCACCTGGAGGTGTACAGCTGCCACCCACCACGGAGCCCGGCCAAGAG 1220
|| ||| | ||||| ||||| ||||| ||||| ||||| |||||

murMU-1  TCAGCCTTACATCTGT.....CATTGTATCCAGCCAAGGA 1374

huMU-1  GCTGCAGCTCACGGAGCTACAAGAACCAGCAGAGCTGGTGGAGTCTGAGC 1270
| ||| | ||||| ||||| ||||| ||||| ||||| ||||| |||||

murMU-1  GAAGAAGTTCGCGGGCTGCCGGGTCTGGAAGAGCAACTGGAGTGTGATG 1424

huMU-1  GTGTGCCCAAGCCAGCTTCTGG.....CCGACAGCCAGAACTCG 1311
| ||| | ||||| ||||| ||||| ||||| ||||| ||||| |||||

murMU-1  GAATGTCTGAGCCTGGTCACTGGTGCATAATCCCCTGGCAGCTGGCCAA 1474

huMU-1  GGGGGCTCAGCTTACAGTGAGGAGAGGGATCGGCCATACGGCCTGGTGTC 1361
| || ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

murMU-1  GCGGTCTCAGCCTACAGTGAGGAGAGAGACCGGCCATATGGTCTGGTGTC 1524

huMU-1  CATTGACACAGTGACTGTGCTAGATGCAGAGGGGCCATGCACCTGGCCCT 1411
||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

murMU-1  CATTGACACAGTGACTGTGGGAGATGCAGAGGGCCTGTGTGTCTGGCCCT 1574

huMU-1  GCAGCTGTGAGGATGACTGGCTACCCAGCCTGGACCTGGATGCTGGCCTG 1461
| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

murMU-1  GTAGCTGTGAGGATGATGGCTATCCAGCCATGAACCTGGATGCTGGCCGA 1624

huMU-1  GAGCCAGCCAGGCCCTAGAGGACCCACTCTTGGATGCAGGGACCACAGT 1511
||| | ||||| ||||| ||||| ||||| ||||| ||||| |||||

murMU-1  GAGTCTGGCCCTAATTCAGAGGATCTGCTCTTGGTACAGACCTGCTTT 1674

huMU-1  CCTGTCTGTGGCTGTGTCTCAGCTGGCAGCCCTGGGCTAGGAGGGCCCC 1561
||||| || ||||| ||||| ||||| ||||| ||||| ||||| |||||

murMU-1  TCTGTCTTGGGGCTGTGTCTCAGGTAGTGGTCTCAGGCTTGGAGGCTCCC 1724

huMU-1  TGGGAAGCCTCCTGGACAGACTAAAGCCACCCCTTGACATGGGGAGGAC 1611
|| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

murMU-1  CAGGCAGCCTACTGGACAGGTTGAGGCTGTCAATTTGCAAAGGAAGGGGAC 1774

huMU-1  TGGGCTGGGGGACTGCCCTGGGGTGGGGGTCACTGGAGGGGTCTCAGA 1661

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FIG.3 (continued)

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murMU-1  ||| || | | | |||| | ||| || ||||| ||| ||
TGGACAGCAGACCCAACCTGGAGAACTGGGTCCCCAGGAGGGGGCTCTGA 1824

huMU-1  GAGTGAGCGGGGCTCACCCCTGGCGGCCTGGATATGGACAGTTTGACA 1711
||||| || || || ||| | || ||||| ||||| |||||

murMU-1  GAGTGAAGCAGGTTCCCCC...CTGGTCTGGACATGGACACATTTGACA 1871

huMU-1  GTGGCTTTGTGGGCTCTGACTGCAGCAGCCCTGTGGAGTGTGACTTCACC 1761
||||||| || || ||||| ||||| ||||| |

murMU-1  GTGGCTTTGCAGGTTCAGACTGTGGCAGCCCCGTGGAGACT..... 1912

huMU-1  AGCCCCGGGGACGAAGGACCCCCCGGAGCTACCTCCGCCAGTGGGTGGT 1811
|| ||||| || || ||||| ||||| ||||| |||

murMU-1  .....GATGAAGGACCCCTCGAAGCTATCTCCGCCAGTGGGTGGT 1953

huMU-1  CATTCCTCCGCCACTTTCGAGCCCTGGACCCAGGCCAGCTAATGAGGCT 1861
|| || || ||| | | || ||||| |||||

murMU-1  CAGGACCCCTCCACCTGTGGACAGTGGAGCCAGAGCAGCTA..... 1995

huMU-1  GACTGGATGTCCAGAGCTGGCCAGGCCACTGGGCTCTGAGCCAGAGACAA 1911
| | | ||| | | || |||||

murMU-1  .GCATATAATAACCAGCTATAGTGAGAAGAGGCCTCTGAGCC..... 2036

huMU-1  TGGGCTTTGAGCCTGATGTTTACAGTGTCTGTGTGTGTGTGTCATATG 2011
||| || || ||| | || ||||| ||||| || ||

murMU-1  TGGCATTACAGTGTGAACATGTAGGGGTGTGTGTGTGTGTGTGTGTG 2086

huMU-1  TGTGTGTGTGCATATGCATGTGTGTGTGTGTGTGTGTGTCTTACTGGACTCA 2061
||||||| || ||||| ||||| ||||| || ||

murMU-1  TGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTGTCTT.GGGTTGTGT 2135

huMU-1  CGGAGCTCACCCATGTGCACAAGTGTGCACAGTAACGTGTTTGTGGTCA 2111
||| || ||||| | || || ||| |

murMU-1  GTTAGCACATCCATGTTGGGATTG.....GTCTGTGTGCTA 2171

huMU-1  ACAGATGACAACAGCCGTCCTCCCTCCTAGGGTCTGTGTGTGCAAGTTGG 2161
|| | | ||| || || || || || || || ||

murMU-1  TGTATTGTAATGCTAAATTCTCTACCCAAAGTTCTAGGCCTACGAGTGAA 2221

huMU-1  TCCACAGCATCTCCGGGGCTTTGTGGGATCAGGGCATTGCCTGTGACTGA 2211
|| || | | || || || || || || || ||

murMU-1  TTCTCATGTTTACAAACTTGCTGTGTAAACCTTG...TTCCTTAATTAA 2268

huMU-1  GCGGAGCCAGCCCTCCAGCGTCTGCCTCCAGGAGCTGCAAGAAGTCCA 2261
| | | ||| || || || || || || || ||

murMU-1  TACCATTGGTTAAATAAAATTGGCTGCAACCAATTACTGGAGGGATTAGA 2318

huMU-1  TATTG.....TTCCTTATCACCTGCCAACAGGAAGCGAAAGGGGATGGAG 2306
|| | || ||||| || || || || || || ||

murMU-1  GGTAGGGGGCTTTTGAGTTACCTGTTTGGAGATGGAGAAGGAGAGAGGAG 2368

huMU-1  TGAGCCCATGGTGACCTCGGGAATGGCAATTTTGGGGCGGCCCTGGAC 2356
|| | || || || || || || || || || ||

murMU-1  AGACCAAGAGGAGAAGGAGGAAGGAGAGGAGAGGAGAGGAGAGGAGGA 2418

huMU-1  GAAGGTCTGAATCCCGACTCTGATACCTTCTGGCTGTGCTACCTGAGCCA 2406
|| | || || || || || || || || || ||

murMU-1  GAGGAGAGGAGAGGAGA.GGAGAGGAGAGGAGAGGCTGCCGTGAGGGGAG 2467

huMU-1  AGTCGCCTCCCTCTCTGGGCTAGAGTTTCCTTATCCAGACAGTGGGGAA 2456
|| || || || || || || || || || || ||

murMU-1  AGGGACCATGAGCCTGTGGCCAGGAGAAACAGCA.....AGTA 2505

huMU-1  GGCATGACACACCTGGGGGAAATTGGCGATGTCACCCGTGTACGGTACGC 2506
| |||| || || ||||| || || || || || ||

murMU-1  TCTGGGGTACACTGGTGAGGAGGTGGCCAGGCCAGC..AGTTAGAAGAGT 2553

huMU-1  AGCCCAGAGCAGACCCTCAATAAAGGTCAGCTTCCTTCAAAAAAAAAA 2556

```


FIG.3 (continued)

murMU-1 || || | |||| || || |||| | |||| || ||||
AGATTAGGGGTGACCTCCAGTATTGTCAAAGCCAATTAAAATAACAAA 2503

huMU-1 AAAAAATCTAGA..... 2567
 |||| | |

murMU-1 AAAAAAAAAAGCGGCGCTCTAGA 2628

FIG.4

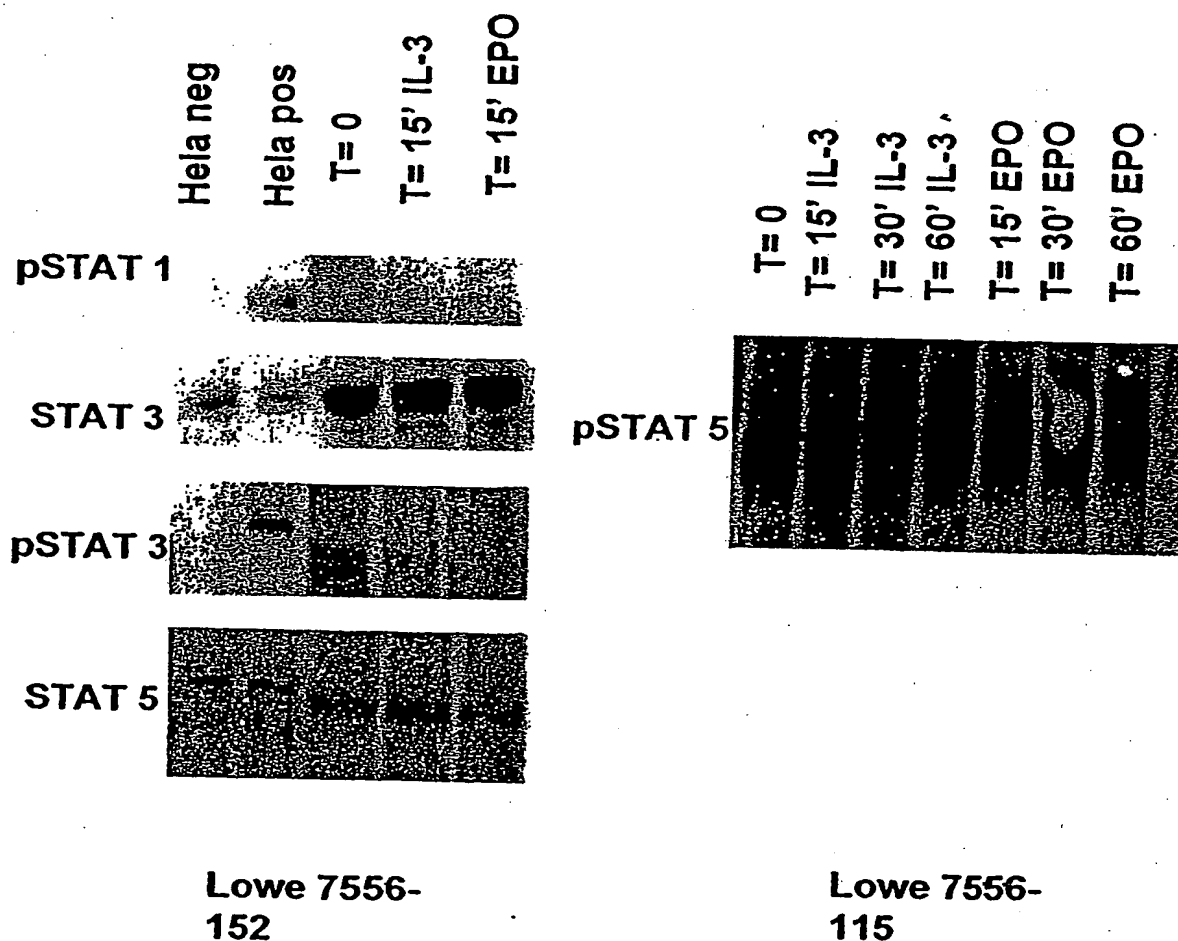
Human MU-1 MPRGWAAPLLLLLLQGGWGCPDLVCYTDYLTQTVICILEMWNLHPSTLTLT 50
 |||| | ||||:| | | || ||||| |: |:| | . || |.||
 MurineMU-1 MPRGPVAALLLLILHGAWSCLDLCYTDYLTITCVLETRSFNPSILSLT 50
 Human MU-1 WQDQYEELKDEATSCSLHRSAHNATHATYTCMDVVFHFMADDIFSVNITD 100
 |||:|||| |: | ||||| || || ||||| . |: |:|:| ||:|
 MurineMU-1 WQDEYEELQDQETFCSLHRSGHNTTHIWYTCMRLSQFLSDEVFIVNVD 100
 Human MU-1 QSGNYSQECGSFLLAESIKPAPPFNVTVTFSGOYNISWRSDYEDPAFYML 150
 |||| |||||..||||||| |||| |||.|| | |:|. |.
 MurineMU-1 QSGNNSQECGSFVLAESIKPAPPLNVTVAFSGRYDISWDSAYDEPSNYVL 150
 Human MU-1 KGKLYEELQYRNRGDPWAVSPRRKLISVDSRSVSLPLEFRKDSSYELQV 200
 :||||||| ||:| | |||||..|||| || ||||:|
 MurineMU-1 RGKLYEELQYRNLRDPYAVRPVTKLISVDSRNVSLPEEFHKDSSYQLQV 200
 Human MU-1 RAGPMPGSSYQGTWSEWSDPVIFQTQSEELKEGWNPHLLLLLLLVIVFIP 250
 || | ||. |:| ||||| ||||| . | . ||. ||:| ||| :
 MurineMU-1 RAAPQPGTSFRGTWSEWSDPVIFQTQAGEPEAGWDPHMLLLAVLIIVL. 249
 Human MU-1 AFWSLKTHPLWRLWKKIWA.VPSPERFFMPLYKGCSDGFKKWVGAPFTGS 299
 | | | ||||| ||. || || |||: ||. |||| |
 MurineMU-1 VFMGLKIHLPWRLWKKIWAPVPTPESFFQPLYREHSGNFKKWVNTPTAS 299
 Human MU-1 SLELGPWSPEVPSTLEVYSCHPPRSPAKRLQLTELQEPALVESDGVPKP 349
 |:| | | | | || | || | | | | | | | | | |
 MurineMU-1 SIELVPQSSTTTTAL.....HLSLYPAKEKKFPGLPGLLEEQLCDGMSEP 344
 Human MU-1 SFW...PTAONSGGSAYSEERDRPYGLVSIIDTVTVLDAEGPCTWPCSCD 396
 | | | ||||| ||||| ||||| ||||| ||||| |||||
 MurineMU-1 GHWCIIPLAAGQAVSAYSEERDRPYGLVSIIDTVTVGDAEGLCVWPCSCD 394
 Human MU-1 DGYPALDLDAGLEPSPGLEDPLLDAGTTVLSCGCVSAGSPGLGGPLGSLL 446
 ||||:|||| | | || || ||||| ||||| |||||
 MurineMU-1 DGYPAMNLDAGRESGPNSDILLVTDPAFLSCGCVSGSGLRLGGSPGSLL 444
 Human MU-1 DRLKPFLADGEDWAGGLPWGGRSPGGVSESEAGSPLAGLMDTDFDSGFVG 496
 |||: | | | | || ||||| ||||| ||||| |||||
 MurineMU-1 DRLRLSFAKEGOWTADPTWRTGSPGGGSESEAGSP.PGLMDTDFDSGFAG 493
 Human MU-1 SDCSSPVECDFTSPGDEGPSPRSYLQWVV.IPPPLSSPGQAS* 539
 ||| |||| ||||| ||||| |||. | | | |
 MurineMU-1 SDCGSPVET.....DEGPSPRSYLQWVVRTPPPVDS.GAQSS. 529

FIG.5

	1		50
humu	---MPRGWAA PLLLLL..LQ GGWG.....	CPDLVCYTDY LQTVICILEM	
mousemu	---MPRGFVA ALLLLI..LH GAWG.....	CLDLTCYTDY LWTITCVLET	
humil2rbc	MAAPALSWRL PLLLLLPLA TSWASAAVNG	TSQFTCFYNS RANISCVWSQ	
	51		100
humu	WN..LHPSTL TLTWQDQYEE LKDEATSCSL	HRSAHNATHA TYTCHM....	
mousemu	RS..PNPSIL SLTWQDEYEE LQDQETFCSL	HRSQHNTTHI WYTCHM....	
humil2rbc	DGALQDTSCQ VHAWPDR....RRWNQTCELLPVSQA SWACNLIGA	
	101		150
humu	.DVHFHMADD IFSVNITDQS GN..YSQECG	SFLLAESIKP APPFNVTVTF	
mousemu	.RLSQFLSDE VFIVNVTDQS GN..NSQECG	SFVLAESIKP APPLNVTVAF	
humil2rbc	PDSQKLTTVD IVTLRVLCRE GVRWRVMAIQ	DFKPFENLRL MAPISLQVH	
	151		200
humu	..SGQYNISW RSDYEDPAFY MLKGKLQYEL	QYRNRGDPWA VSPRRKLISV	
mousemu	..SGRYDISW DSAYDEPSNY VLRGKLQYEL	QYRNLRDPA VRPVTKLISV	
humil2rbc	VETHRCNISW E..ISQASHY FER.HLEFEA	RTLSPGHTWE EAP...LTL	
	201		250
humu	DSRSVSLPL EFRKDSSYEL QVRAGPMPS	SYQGTWSEWS DPVIFQTQS.	
mousemu	DSRNVSLLPE EFHKDSSYQL QVRAAPQPGT	SFRGTWSEWS DPVIFQTQA.	
humil2rbc	KQKQEWICLE TLTPDTQYEF QVRVKPLQGE	F..TTWSPWS QPLAFRTKPA	
	251		300
humu	..EELKEGWN PHLLLLL...LLVIVFIPAF	WSLKTHPLWR LWKKINA.VP	
mousemu	..GEPEAGWD PHMLLLL...AVLIIVL.VF	MGLKIHLPWR LWKKINAVP	
humil2rbc	ALGKDTIPWL GHLLVGLSGA FGFIILVYLL	INCRNTGPW. LKKVLKCNTP	
	301		350
humu	<u>SPERFFMPLY</u> KGCSGDFKKW VGAPFTGSSL	ELGPWSPEVP <u>STLEVYSCHP</u>	
mousemu	<u>TESFFQPLY</u> REHSGNFKKW VNTPFTASSI	ELVQSSSTT <u>SAL.....HL</u>	
humil2rbc	DPSKFSSQLS SEHGSDVQKW LSSPFPSSSF	SPGGLAPEIS PLEVLERDKV	
	351		400
humu	<u>PRSPAKRLQL</u> TELQEPAA..E LVESDGVPKP	SFW...PTAQ NSGGSAYSEE	
mousemu	<u>SLYPAKEKKF</u> PGLPGLA..E QLECDGMSEP	GHWCIIPLAA GQAVSAYSEE	
humil2rbc	TQLLLOQDKV PEPASLSSNH SLTSCFTNQG	YFFFHLPDAL EIEACQVYFT	
	401		450
humu	RDRPYGLVSI DTVTVLDAEG PC...TWPCS	CEDDGYPALD LDAGLEPSPG	
mousemu	RDRPYGLVSI DTVTVGDAEG LC...VWPCS	CEDDGYPAMN LDAGRESGPN	
humil2rbc	YD.PYSEEDP DEGVAGAPTG SSPQPLQPLS	GEDDAYCTF.PS	
	451		500
humu	LEDPLLDAGT TVLSCGCVSA GSPGLGGPLG	SLLDRLKPPL AD..GEDWAG	
mousemu	SEDLLLVTDP AFLSCGCVSG SGLRLGGSPG	SLLDRLRLSF AK..EGDWTA	
humil2rbc	RDDLLLF.S.P SLL..GGPSP PSTAPGGS.G	AGEERMPPSL QERVPRWDWP	
	501		550
humu	GLPWGGRSPG GVSESEAGSP LAGLMDTDF	SGFVGSDCSS PVECDFTSPG	
mousemu	DPTWRTGSPG GGSESEAGSP .PGLMDTDF	SGFAGSDCGS PVET.....	
humil2rbc	Q.PLGPPTPG VPDLVDFQPP P...ELVLRE	AGEEVPDAG. PRE.GVSFPW	
	551		588
humu	DEGPPRSYLR QWVVIPPLS SPGPQAS+---	-----	
mousemu	DEGPPRSYLR QWVVRTFPFV DSGAQSS---	-----	
humil2rbc	SRPPGQGEFR ALNARLPLNT DAYLSLQELQ	GQDPHTLV	

FIG.6

Signaling through MU-1



atg Met 1	aaa Lys	ttc Phe	tta Leu	gtc Val 5	aac Asn	gtt Val	gcc Ala	ctt Leu	gtt Val 10	ttt Phe	atg Met	gtc Val	gtg Val	tac Tyr 15	att Ile	48
tct Ser	tac Tyr	atc Ile	tat Tyr 20	gcc Ala	ggc Gly	agc Ser	gga Gly	cac His 25	cac His	cat His	cat His	cac His	cac His 30	ggg Gly	agc Ser	96
ggc Gly	gac Asp	tat Tyr 35	aaa Lys	gac Asp	gat Asp	gac Asp	gat Asp 40	aag Lys	ggg Gly	tcc Ser	gga Gly	tgc Cys 45	ccc Pro	gac Asp	ctc Leu	144
gtc Val	tgc Cys 50	tac Tyr	acc Thr	gat Asp	tac Tyr	ctc Leu 55	cag Gln	acg Thr	gtc Val	atc Ile	tgc Cys 60	atc Ile	ctg Leu	gaa Glu	atg Met	192
tgg Trp 65	aac Asn	ctc Leu	cac His	ccc Pro	agc Ser 70	acg Thr	ctc Leu	acc Thr	ctt Leu	acc Thr 75	tgg Trp	caa Gln	gac Asp	cag Gln	tat Tyr 80	240
gaa Glu	gag Glu	ctg Leu	aag Lys	gac Asp 85	gag Glu	gcc Ala	acc Thr	tcc Ser	tgc Cys 90	agc Ser	ctc Leu	cac His	agg Arg	tcg Ser 95	gcc Ala	288
cac His	aat Asn	gcc Ala	acg Thr 100	cat His	gcc Ala	acc Thr	tac Tyr	acc Thr 105	tgc Cys	cac His	atg Met	gat Asp	gta Val 110	ttc Phe	cac His	336
ttc Phe	atg Met	gcc Ala 115	gac Asp	gac Asp	att Ile	ttc Phe	agt Ser 120	gtc Val	aac Asn	atc Ile	aca Thr	gac Asp 125	cag Gln	tct Ser	ggc Gly	384
aac Asn	tac Tyr 130	tcc Ser	cag Gln	gag Glu	tgt Cys	ggc Gly 135	agc Ser	ttt Phe	ctc Leu	ctg Leu	gct Ala 140	gag Glu	agc Ser	atc Ile	aag Lys	432
ccg Pro 145	gct Ala	ccc Pro	cct Pro	ttc Phe	aac Asn 150	gtg Val	act Thr	gtg Val	acc Thr	ttc Phe 155	tca Ser	gga Gly	cag Gln	tat Tyr	aat Asn 160	480
atc Ile	tcc Ser	tgg Trp	cgc Arg	tca Ser 165	gat Asp	tac Tyr	gaa Glu	gac Asp	cct Pro 170	gcc Ala	ttc Phe	tac Tyr	atg Met	ctg Leu 175	aag Lys	528
ggc Gly	aag Lys	ctt Leu	cag Gln 180	tat Tyr	gag Glu	ctg Leu	cag Gln	tac Tyr 185	agg Arg	aac Asn	cgg Arg	gga Gly	gac Asp 190	ccc Pro	tgg Trp	576
gct Ala	gtg Val	agt Ser 195	ccg Pro	agg Arg	aga Arg	aag Lys	ctg Leu 200	atc Ile	tca Ser	gtg Val	gac Asp 205	tca Ser	aga Arg	agt Ser	gtc Val	624
tcc Ser	ctc Leu 210	ctc Leu	ccc Pro	ctg Leu	gag Glu	ttc Phe 215	cgc Arg	aaa Lys	gac Asp	tcg Ser	agc Ser 220	tat Tyr	gag Glu	ctg Leu	cag Gln	672

Figure 7A

gtg	cgg	gca	ggg	ccc	atg	cct	ggc	tcc	tcc	tac	cag	ggg	acc	tgg	agt		720
Val	Arg	Ala	Gly	Pro	Met	Pro	Gly	Ser	Ser	Tyr	Gln	Gly	Thr	Trp	Ser		
225					230					235					240		
gaa	tgg	agt	gac	ccg	gtc	atc	ttt	cag	acc	cag	tca	gag	gag	tta	aag		768
Glu	Trp	Ser	Asp	Pro	Val	Ile	Phe	Gln	Thr	Gln	Ser	Glu	Glu	Leu	Lys		
				245					250					255			
gaa	ggc	tgg	aac	taa	tga	SEQ ID NO: 22											786
Glu	Gly	Trp	Asn	SEQ ID NO: 23													
	260																

Figure 7B

gcggccgcac cacc	atg Met 1	ccg Pro	cgt Arg	ggc Gly	tgg Trp 5	gcc Ala	gcc Ala	ccc Pro	ttg Leu	ctc Leu 10	ctg Leu	ctg Leu	50
ctg ctc cag gga ggc tgg ggc tgc ccc gac ctc gtc tgc tac acc gat	Leu Leu Gln Gly Gly Trp Gly Cys Pro Asp Leu Val Cys Tyr Thr Asp	15			20				25				98
tac ctc cag acg gtc atc tgc atc ctg gaa atg tgg aac ctc cac ccc	Tyr Leu Gln Thr Val Ile Cys Ile Leu Glu Met Trp Asn Leu His Pro	30			35			40					146
agc acg ctc acc ctt acc tgg caa gac cag tat gaa gag ctg aag gac	Ser Thr Leu Thr Leu Thr Trp Gln Asp Gln Tyr Glu Glu Leu Lys Asp	45			50			55				60	194
gag gcc acc tcc tgc agc ctc cac agg tgc gcc cac aat gcc acg cat	Glu Ala Thr Ser Cys Ser Leu His Arg Ser Ala His Asn Ala Thr His			65			70				75		242
gcc acc tac acc tgc cac atg gat gta ttc cac ttc atg gcc gac gac	Ala Thr Tyr Thr Cys His Met Asp Val Phe His Phe Met Ala Asp Asp	80				85				90			290
att ttc agt gtc aac atc aca gac cag tct ggc aac tac tcc cag gag	Ile Phe Ser Val Asn Ile Thr Asp Gln Ser Gly Asn Tyr Ser Gln Glu	95				100				105			338
tgt ggc agc ttt ctc ctg gct gag agc atc aag ccg gct ccc cct ttc	Cys Gly Ser Phe Leu Leu Ala Glu Ser Ile Lys Pro Ala Pro Pro Phe	110			115				120				386
aac gtg act gtg acc ttc tca gga cag tat aat atc tcc tgg cgc tca	Asn Val Thr Val Thr Phe Ser Gly Gln Tyr Asn Ile Ser Trp Arg Ser	125			130			135				140	434
gat tac gaa gac cct gcc ttc tac atg ctg aag ggc aag ctt cag tat	Asp Tyr Glu Asp Pro Ala Phe Tyr Met Leu Lys Gly Lys Leu Gln Tyr			145				150				155	482
gag ctg cag tac agg aac cgg gga gac ccc tgg gct gtg agt ccg agg	Glu Leu Gln Tyr Arg Asn Arg Gly Asp Pro Trp Ala Val Ser Pro Arg			160			165				170		530
aga aag ctg atc tca gtg gac tca aga agt gtc tcc ctc ctc ccc ctg	Arg Lys Leu Ile Ser Val Asp Ser Arg Ser Val Ser Leu Leu Pro Leu	175				180				185			578
gag ttc cgc aaa gac tgc agc tat gag ctg cag gtg cgg gca ggg ccc	Glu Phe Arg Lys Asp Ser Ser Tyr Glu Leu Gln Val Arg Ala Gly Pro	190				195				200			626
atg cct ggc tcc tcc tac cag ggg acc tgg agt gaa tgg agt gac ccg	Met Pro Gly Ser Ser Tyr Gln Gly Thr Trp Ser Glu Trp Ser Asp Pro	205			210				215			220	674

Figure 8A

gtc Val	atc Ile	ttt Phe	cag Gln	acc Thr 225	cag Gln	tca Ser	gag Glu	gag Glu	tta Leu 230	aag Lys	gaa Glu	ggc Gly	tgg Trp	aac Asn 235	ggc Gly	722
tcc Ser	ggc Gly	tct Ser	aga Arg 240	gac Asp	aaa Lys	act Thr	cac His	aca Thr 245	tgc Cys	cca Pro	ccg Pro	tgc Cys	cca Pro 250	gca Ala	cct Pro	770
gaa Glu	ctc Leu	ctg Leu 255	ggg Gly	gga Gly	ccg Pro	tca Ser	gtc Val 260	ttc Phe	ctc Leu	ttc Phe	ccc Pro	cca Pro 265	aaa Lys	ccc Pro	aag Lys	818
gac Asp	acc Thr 270	ctc Leu	atg Met	atc Ile	tcc Ser	cgg Arg 275	acc Thr	cct Pro	gag Glu	gtc Val	aca Thr 280	tgc Cys	gtg Val	gtg Val	gtg Val	866
gac Asp 285	gtg Val	agc Ser	cac His	gaa Glu	gac Asp 290	cct Pro	gag Glu	gtc Val	aag Lys	ttc Phe 295	aac Asn	tgg Trp	tac Tyr	gtg Val	gac Asp 300	914
ggc Gly	gtg Val	gag Glu	gtg Val	cat His 305	aat Asn	gcc Ala	aag Lys	aca Thr	aag Lys 310	ccg Pro	cgg Arg	gag Glu	gag Glu	cag Gln 315	tac Tyr	962
aac Asn	agc Ser	acg Thr	tac Tyr 320	cgt Arg	gtg Val	gtc Val	agc Ser	gtc Val 325	ctc Leu	acc Thr	gtc Val	ctg Leu	cac His 330	cag Gln	gac Asp	1010
tgg Trp	ctg Leu	aat Asn 335	ggc Gly	aag Lys	gag Glu	tac Tyr	aag Lys 340	tgc Cys	aag Lys	gtc Val	tcc Ser	aac Asn 345	aaa Lys	gcc Ala	ctc Leu	1058
cca Pro	gtc Val 350	ccc Pro	atc Ile	gag Glu	aaa Lys	acc Thr 355	atc Ile	tcc Ser	aaa Lys	gcc Ala	aaa Lys 360	ggg Gly	cag Gln	ccc Pro	cga Arg	1106
gaa Glu 365	cca Pro	cag Gln	gtg Val	tac Tyr	acc Thr 370	ctg Leu	ccc Pro	cca Pro	tcc Ser	cgg Arg 375	gag Glu	gag Glu	atg Met	acc Thr	aag Lys 380	1154
aac Asn	cag Gln	gtc Val	agc Ser	ctg Leu 385	acc Thr	tgc Cys	ctg Leu	gtc Val	aaa Lys 390	ggc Gly	ttc Phe	tat Tyr	ccc Pro	agc Ser 395	gac Asp	1202
atc Ile	gcc Ala	gtg Val	gag Glu 400	tgg Trp	gag Glu	agc Ser	aat Asn	ggg Gly 405	cag Gln	ccg Pro	gag Glu	aac Asn	aac Asn 410	tac Tyr	aag Lys	1250
acc Thr	acg Thr	cct Pro 415	ccc Pro	gtg Val	ctg Leu	gac Asp	tcc Ser 420	gac Asp	ggc Gly	tcc Ser	ttc Phe	ttc Phe 425	ctc Leu	tat Tyr	agc Ser	1298
aag Lys	ctc Leu 430	acc Thr	gtg Val	gac Asp	aag Lys	agc Ser 435	agg Arg	tgg Trp	cag Gln	cag Gln	ggg Gly 440	aac Asn	gtc Val	ttc Phe	tca Ser	1346

Figure 8B

tgc	tcc	gtg	atg	cat	gag	gct	ctg	cac	aac	cac	tac	acg	cag	aag	agc	1394
Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr	Thr	Gln	Lys	Ser	
445					450					455					460	
ctc	tcc	ctg	tcc	ccg	ggt	aaa	tgagtgaatt	c	SEQ ID NO: 24							1426
Leu	Ser	Leu	Ser	Pro	Gly	Lys	SEQ ID NO: 25									
				465												

Figure 8C

Q ID NO:26	gcggccgcac	cacc	atg	ccg	cgt	ggc	tgg	gcc	gcc	ccc	ttg	ctc	ctg	ctg	50		
Q ID NO:27			Met	Pro	Arg	Gly	Trp	Ala	Ala	Pro	Leu	Leu	Leu	Leu			
			1				5				10						
	ctg	ctc	cag	gga	ggc	tgg	ggc	tgc	ccc	gac	ctc	gtc	tgc	tac	acc	gat	98
	Leu	Leu	Gln	Gly	Gly	Trp	Gly	Cys	Pro	Asp	Leu	Val	Cys	Tyr	Thr	Asp	
			15				20					25					
	tac	ctc	cag	acg	gtc	atc	tgc	atc	ctg	gaa	atg	tgg	aac	ctc	cac	ccc	146
	Tyr	Leu	Gln	Thr	Val	Ile	Cys	Ile	Leu	Glu	Met	Trp	Asn	Leu	His	Pro	
		30					35					40					
	agc	acg	ctc	acc	ctt	acc	tgg	caa	gac	cag	tat	gaa	gag	ctg	aag	gac	194
	Ser	Thr	Leu	Thr	Leu	Thr	Trp	Gln	Asp	Gln	Tyr	Glu	Glu	Leu	Lys	Asp	
	45					50					55					60	
	gag	gcc	acc	tcc	tgc	agc	ctc	cac	agg	tcg	gcc	cac	aat	gcc	acg	cat	242
	Glu	Ala	Thr	Ser	Cys	Ser	Leu	His	Arg	Ser	Ala	His	Asn	Ala	Thr	His	
					65				70						75		
	gcc	acc	tac	acc	tgc	cac	atg	gat	gta	ttc	cac	ttc	atg	gcc	gac	gac	290
	Ala	Thr	Tyr	Thr	Cys	His	Met	Asp	Val	Phe	His	Phe	Met	Ala	Asp	Asp	
				80					85					90			
	att	ttc	agt	gtc	aac	atc	aca	gac	cag	tct	ggc	aac	tac	tcc	cag	gag	338
	Ile	Phe	Ser	Val	Asn	Ile	Thr	Asp	Gln	Ser	Gly	Asn	Tyr	Ser	Gln	Glu	
			95					100					105				
	tgt	ggc	agc	ttt	ctc	ctg	gct	gag	agc	atc	aag	ccg	gct	ccc	cct	ttc	386
	Cys	Gly	Ser	Phe	Leu	Leu	Ala	Glu	Ser	Ile	Lys	Pro	Ala	Pro	Pro	Phe	
		110					115					120					
	aac	gtg	act	gtg	acc	ttc	tca	gga	cag	tat	aat	atc	tcc	tgg	cgc	tca	434
	Asn	Val	Thr	Val	Thr	Phe	Ser	Gly	Gln	Tyr	Asn	Ile	Ser	Trp	Arg	Ser	
	125					130					135					140	
	gat	tac	gaa	gac	cct	gcc	ttc	tac	atg	ctg	aag	ggc	aag	ctt	cag	tat	482
	Asp	Tyr	Glu	Asp	Pro	Ala	Phe	Tyr	Met	Leu	Lys	Gly	Lys	Leu	Gln	Tyr	
					145					150					155		
	gag	ctg	cag	tac	agg	aac	cgg	gga	gac	ccc	tgg	gct	gtg	agt	ccg	agg	530
	Glu	Leu	Gln	Tyr	Arg	Asn	Arg	Gly	Asp	Pro	Trp	Ala	Val	Ser	Pro	Arg	
				160					165					170			
	aga	aag	ctg	atc	tca	gtg	gac	tca	aga	agt	gtc	tcc	ctc	ctc	ccc	ctg	578
	Arg	Lys	Leu	Ile	Ser	Val	Asp	Ser	Arg	Ser	Val	Ser	Leu	Leu	Pro	Leu	
			175					180					185				
	gag	ttc	cgc	aaa	gac	tcg	agc	tat	gag	ctg	cag	gtg	cgg	gca	ggg	ccc	626
	Glu	Phe	Arg	Lys	Asp	Ser	Ser	Tyr	Glu	Leu	Gln	Val	Arg	Ala	Gly	Pro	
		190					195					200					
	atg	cct	ggc	tcc	tcc	tac	cag	ggg	acc	tgg	agt	gaa	tgg	agt	gac	ccg	674
	Met	Pro	Gly	Ser	Ser	Tyr	Gln	Gly	Thr	Trp	Ser	Glu	Trp	Ser	Asp	Pro	
	205					210					215					220	

Figure 9A

gtc Val	atc Ile	ttt Phe	cag Gln	acc Thr 225	cag Gln	tca Ser	gag Glu	gag Glu	tta Leu 230	aag Lys	gaa Glu	ggc Gly	tgg Trp	aac Asn 235	ggc Gly	722
tcc Ser	ggc Gly	tct Ser	aga Arg 240	gac Asp	aaa Lys	act Thr	cac His	aca Thr 245	tgc Cys	cca Pro	ccg Pro	tgc Cys	cca Pro 250	gca Ala	cct Pro	770
gaa Glu	ctc Leu	ctg Leu 255	ggg Gly	gga Gly	ccg Pro	tca Ser	gtc Val 260	ttc Phe	ctc Leu	ttc Phe	ccc Pro	cca Pro 265	aaa Lys	ccc Pro	aag Lys	818
gac Asp	acc Thr 270	ctc Leu	atg Met	atc Ile	tcc Ser	cgg Arg 275	acc Thr	cct Pro	gag Glu	gtc Val	aca Thr 280	tgc Cys	gtg Val	gtg Val	gtg Val	866
gac Asp 285	gtg Val	agc Ser	cac His	gaa Glu	gac Asp 290	cct Pro	gag Glu	gtc Val	aag Lys	ttc Phe 295	aac Asn	tgg Trp	tac Tyr	gtg Val	gac Asp 300	914
ggc Gly	gtg Val	gag Glu	gtg Val	cat His 305	aat Asn	gcc Ala	aag Lys	aca Thr	aag Lys 310	ccg Pro	cgg Arg	gag Glu	gag Glu	cag Gln 315	tac Tyr	962
aac Asn	agc Ser	acg Thr	tac Tyr 320	cgt Arg	gtg Val	gtc Val	agc Ser	gtc Val 325	ctc Leu	acc Thr	gtc Val	ctg Leu	cac His 330	cag Gln	gac Asp	1010
tgg Trp	ctg Leu	aat Asn 335	ggc Gly	aag Lys	gag Glu	tac Tyr	aag Lys 340	tgc Cys	aag Lys	gtc Val	tcc Ser	aac Asn 345	aaa Lys	gcc Ala	ctc Leu	1058
cca Pro	gtc Val 350	ccc Pro	atc Ile	gag Glu	aaa Lys	acc Thr 355	atc Ile	tcc Ser	aaa Lys	gcc Ala 360	aaa Lys	ggg Gly	cag Gln	ccc Pro	cga Arg	1106
gaa Glu 365	cca Pro	cag Gln	gtg Val	tac Tyr	acc Thr 370	ctg Leu	ccc Pro	cca Pro	tcc Ser	cgg Arg 375	gag Glu	gag Glu	atg Met	acc Thr	aag Lys 380	1154
aac Asn	cag Gln	gtc Val	agc Ser	ctg Leu 385	acc Thr	tgc Cys	ctg Leu	gtc Val	aaa Lys 390	ggc Gly	ttc Phe	tat Tyr	ccc Pro	agc Ser 395	gac Asp	1202
atc Ile	gcc Ala	gtg Val	gag Glu 400	tgg Trp	gag Glu	agc Ser	aat Asn	ggg Gly 405	cag Gln	ccg Pro	gag Glu	aac Asn	aac Asn 410	tac Tyr	aag Lys	1250
acc Thr	acg Thr	cct Pro 415	ccc Pro	gtg Val	ctg Leu	gac Asp	tcc Ser 420	gac Asp	ggc Gly	tcc Ser	ttc Phe	ttc Phe 425	ctc Leu	tat Tyr	agc Ser	1298
aag Lys	ctc Leu 430	acc Thr	gtg Val	gac Asp	aag Lys	agc Ser 435	agg Arg	tgg Trp	cag Gln	cag Gln	ggg Gly 440	aac Asn	gtc Val	ttc Phe	tca Ser	1346

Figure 9B

tgc	tcc	gtg	atg	cat	gag	gct	ctg	cac	aac	cac	tac	acg	cag	aag	agc		1394
Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr	Thr	Gln	Lys	Ser		
445					450					455					460		
ctc	tcc	ctg	tcc	ccg	ggt	aaa	tca	gga	atg	gca	tca	atg	aca	gga	ggt		1442
Leu	Ser	Leu	Ser	Pro	Gly	Lys	Ser	Gly	Met	Ala	Ser	Met	Thr	Gly	Gly		
				465					470					475			
caa	caa	atg	ggt	tct	gga	tct	cat	cat	cat	cat	cat	cat	tct	gga	ggt		1490
Gln	Gln	Met	Gly	Ser	Gly	Ser	His	His	His	His	His	His	Ser	Gly	Gly		
			480					485					490				
tgagaattc																	1499

Figure 9C

gcggccgcac	cacc	atg	ccg	cgt	ggc	tgg	gcc	gcc	ccc	ttg	ctc	ctg	ctg	50		
		Met	Pro	Arg	Gly	Trp	Ala	Ala	Pro	Leu	Leu	Leu	Leu			
		1				5				10						
ctg	ctc	cag	gga	ggc	tgg	ggc	tgc	ccc	gac	ctc	gtc	tgc	tac	acc	gat	98
Leu	Leu	Gln	Gly	Gly	Trp	Gly	Cys	Pro	Asp	Leu	Val	Cys	Tyr	Thr	Asp	
		15				20						25				
tac	ctc	cag	acg	gtc	atc	tgc	atc	ctg	gaa	atg	tgg	aac	ctc	cac	ccc	146
Tyr	Leu	Gln	Thr	Val	Ile	Cys	Ile	Leu	Glu	Met	Trp	Asn	Leu	His	Pro	
	30					35					40					
agc	acg	ctc	acc	ctt	acc	tgg	caa	gac	cag	tat	gaa	gag	ctg	aag	gac	194
Ser	Thr	Leu	Thr	Leu	Thr	Trp	Gln	Asp	Gln	Tyr	Glu	Glu	Leu	Lys	Asp	
45					50					55					60	
gag	gcc	acc	tcc	tgc	agc	ctc	cac	agg	tgc	gcc	cac	aat	gcc	acg	cat	242
Glu	Ala	Thr	Ser	Cys	Ser	Leu	His	Arg	Ser	Ala	His	Asn	Ala	Thr	His	
				65					70					75		
gcc	acc	tac	acc	tgc	cac	atg	gat	gta	ttc	cac	ttc	atg	gcc	gac	gac	290
Ala	Thr	Tyr	Thr	Cys	His	Met	Asp	Val	Phe	His	Phe	Met	Ala	Asp	Asp	
			80					85					90			
att	ttc	agt	gtc	aac	atc	aca	gac	cag	tct	ggc	aac	tac	tcc	cag	gag	338
Ile	Phe	Ser	Val	Asn	Ile	Thr	Asp	Gln	Ser	Gly	Asn	Tyr	Ser	Gln	Glu	
		95					100					105				
tgt	ggc	agc	ttt	ctc	ctg	gct	gag	agc	atc	aag	ccg	gct	ccc	cct	ttc	386
Cys	Gly	Ser	Phe	Leu	Leu	Ala	Glu	Ser	Ile	Lys	Pro	Ala	Pro	Pro	Phe	
	110					115					120					
aac	gtg	act	gtg	acc	ttc	tca	gga	cag	tat	aat	atc	tcc	tgg	cgc	tca	434
Asn	Val	Thr	Val	Thr	Phe	Ser	Gly	Gln	Tyr	Asn	Ile	Ser	Trp	Arg	Ser	
125					130					135					140	
gat	tac	gaa	gac	cct	gcc	ttc	tac	atg	ctg	aag	ggc	aag	ctt	cag	tat	482
Asp	Tyr	Glu	Asp	Pro	Ala	Phe	Tyr	Met	Leu	Lys	Gly	Lys	Leu	Gln	Tyr	
				145					150					155		
gag	ctg	cag	tac	agg	aac	cgg	gga	gac	ccc	tgg	gct	gtg	agt	ccg	agg	530
Glu	Leu	Gln	Tyr	Arg	Asn	Arg	Gly	Asp	Pro	Trp	Ala	Val	Ser	Pro	Arg	
			160					165					170			
aga	aag	ctg	atc	tca	gtg	gac	tca	aga	agt	gtc	tcc	ctc	ctc	ccc	ctg	578
Arg	Lys	Leu	Ile	Ser	Val	Asp	Ser	Arg	Ser	Val	Ser	Leu	Leu	Pro	Leu	
		175					180					185				
gag	ttc	cgc	aaa	gac	tcg	agc	tat	gag	ctg	cag	gtg	cgg	gca	ggg	ccc	626
Glu	Phe	Arg	Lys	Asp	Ser	Ser	Tyr	Glu	Leu	Gln	Val	Arg	Ala	Gly	Pro	
	190					195					200					
atg	cct	ggc	tcc	tcc	tac	cag	ggg	acc	tgg	agt	gaa	tgg	agt	gac	ccg	674
Met	Pro	Gly	Ser	Ser	Tyr	Gln	Gly	Thr	Trp	Ser	Glu	Trp	Ser	Asp	Pro	
205					210					215					220	

Figure 10A

gtc atc ttt cag acc cag tca gag gag tta aag gaa ggc tgg aac ggc	722
Val Ile Phe Gln Thr Gln Ser Glu Glu Leu Lys Glu Gly Trp Asn Gly	
225	
tcc ggc tct aga gac aaa act cac aca tgc cca ccg tgc cca gca cct	770
Ser Gly Ser Arg Asp Lys Thr His Thr Cys Pro Pro Cys Pro Ala Pro	
240	
245	
250	
gaa gcc ctg ggg gca ccg tca gtc ttc ctc ttc ccc cca aaa ccc aag	818
Glu Ala Leu Gly Ala Pro Ser Val Phe Leu Phe Pro Pro Lys Pro Lys	
255	
260	
265	
gac acc ctc atg atc tcc cgg acc cct gag gtc aca tgc gtg gtg gtg	866
Asp Thr Leu Met Ile Ser Arg Thr Pro Glu Val Thr Cys Val Val Val	
270	
275	
280	
gac gtg agc cac gaa gac cct gag gtc aag ttc aac tgg tac gtg gac	914
Asp Val Ser His Glu Asp Pro Glu Val Lys Phe Asn Trp Tyr Val Asp	
285	
290	
295	
300	
ggc gtg gag gtg cat aat gcc aag aca aag ccg cgg gag gag cag tac	962
Gly Val Glu Val His Asn Ala Lys Thr Lys Pro Arg Glu Glu Gln Tyr	
305	
310	
315	
aac agc acg tac cgt gtg gtc agc gtc ctc acc gtc ctg cac cag gac	1010
Asn Ser Thr Tyr Arg Val Val Ser Val Leu Thr Val Leu His Gln Asp	
320	
325	
330	
tgg ctg aat ggc aag gag tac aag tgc aag gtc tcc aac aaa gcc ctc	1058
Trp Leu Asn Gly Lys Glu Tyr Lys Cys Lys Val Ser Asn Lys Ala Leu	
335	
340	
345	
cca gcc ccc atc gag aaa acc atc tcc aaa gcc aaa ggg cag ccc cga	1106
Pro Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala Lys Gly Gln Pro Arg	
350	
355	
360	
gaa cca cag gtg tac acc ctg ccc cca tcc cgg gag gag atg acc aag	1154
Glu Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg Glu Glu Met Thr Lys	
365	
370	
375	
380	
aac cag gtc agc ctg acc tgc ctg gtc aaa ggc ttc tat ccc agc gac	1202
Asn Gln Val Ser Leu Thr Cys Leu Val Lys Gly Phe Tyr Pro Ser Asp	
385	
390	
395	
atc gcc gtg gag tgg gag agc aat ggg cag ccg gag aac aac tac aag	1250
Ile Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu Asn Asn Tyr Lys	
400	
405	
410	
acc acg cct ccc gtg ctg gac tcc gac ggc tcc ttc ttc ctc tat agc	1298
Thr Thr Pro Val Leu Asp Ser Asp Gly Ser Phe Phe Leu Tyr Ser	
415	
420	
425	
aag ctc acc gtg gac aag agc agg tgg cag cag ggg aac gtc ttc tca	1346
Lys Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly Asn Val Phe Ser	
430	
435	
440	

Figure 10B

tgc	tcc	gtg	atg	cat	gag	gct	ctg	cac	aac	cac	tac	acg	cag	aag	agc	1394
Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr	Thr	Gln	Lys	Ser	
445					450					455					460	
ctc	tcc	ctg	tcc	ccg	ggt	aaa	tgagtgaatt	c	SEQ ID NO: 28							1426
Leu	Ser	Leu	Ser	Pro	Gly	Lys	SEQ ID NO: 29									
				465												

Figure 10C

atg Met 1	ccg Pro	cgt Arg	ggc Gly 5	tgg Trp	gcc Ala	gcc Ala	ccc Pro	ttg Leu 10	ctc Leu	ctg Leu	ctg Leu	ctc Leu	cag Gln 15	gga Gly	48	
ggc Gly	tgg Trp	ggc Gly	tgc Cys 20	ccc Pro	gac Asp	ctc Leu	gtc Val	tgc Cys 25	tac Tyr	acc Thr	gat Asp	tac Tyr	ctc Leu 30	cag Gln	acg Thr	96
gtc Val	atc Ile	tgc Cys 35	atc Ile	ctg Leu	gaa Glu	atg Met	tgg Trp 40	aac Asn	ctc Leu	cac His	ccc Pro	agc Ser 45	acg Thr	ctc Leu	acc Thr	144
ctt Leu	acc Thr 50	tgg Trp	caa Gln	gac Asp	cag Gln	tat Tyr 55	gaa Glu	gag Glu	ctg Leu	aag Lys	gac Asp 60	gag Glu	gcc Ala	acc Thr	tcc Ser	192
tgc Cys 65	agc Ser	ctc Leu	cac His	agg Arg	tcg Ser 70	gcc Ala	cac His	aat Asn	gcc Ala	acg Thr 75	cat His	gcc Ala	acc Thr	tac Tyr	acc Thr 80	240
tgc Cys	cac His	atg Met	gat Asp	gta Val 85	ttc Phe	cac His	ttc Phe	atg Met	gcc Ala 90	gac Asp	gac Asp	att Ile	ttc Phe	agt Ser 95	gtc Val	288
aac Asn	atc Ile	aca Thr	gac Asp 100	cag Gln	tct Ser	ggc Gly	aac Asn	tac Tyr 105	tcc Ser	cag Gln	gag Glu	tgt Cys	ggc Gly 110	agc Ser	ttt Phe	336
ctc Leu	ctg Leu	gct Ala 115	gag Glu	agc Ser	atc Ile	aag Lys	ccg Pro 120	gct Ala	ccc Pro	cct Pro	ttc Phe	aac Asn 125	gtg Val	act Thr	gtg Val	384
acc Thr	ttc Phe 130	tca Ser	gga Gly	cag Gln	tat Tyr	aat Asn 135	atc Ile	tcc Ser	tgg Trp	cgc Arg	tca Ser 140	gat Asp	tac Tyr	gaa Glu	gac Asp	432
cct Pro 145	gcc Ala	ttc Phe	tac Tyr	atg Met	ctg Leu 150	aag Lys	ggc Gly	aag Lys	ctt Leu	cag Gln 155	tat Tyr	gag Glu	ctg Leu	cag Gln	tac Tyr 160	480
agg Arg	aac Asn	cgg Arg	gga Gly	gac Asp 165	ccc Pro	tgg Trp	gct Ala	gtg Val	agt Ser 170	ccg Pro	agg Arg	aga Arg	aag Lys	ctg Leu 175	atc Ile	528
tca Ser	gtg Val	gac Asp	tca Ser 180	aga Arg	agt Ser	gtc Val	tcc Ser 185	ctc Leu 185	ctc Leu	ccc Pro	ctg Leu	gag Glu	ttc Phe 190	cgc Arg	aaa Lys	576
gac Asp	tcg Ser	agc Ser	tat Tyr 195	gag Glu	ctg Leu	cag Gln	gtg Val 200	cgg Arg	gca Ala	ggg Gly	ccc Pro	atg Met 205	cct Pro	ggc Gly	tcc Ser	624
tcc Ser	tac Tyr 210	cag Gln	ggg Gly	acc Thr	tgg Trp	agt Ser 215	gaa Glu	tgg Trp	agt Ser	gac Asp	ccg Pro 220	gtc Val	atc Ile	ttt Phe	cag Gln	672

Figure 11A

acc	cag	tca	gag	gag	tta	aag	gaa	ggc	tg	aac	aaa	acc	gaa	acc	tcc	720
Thr	Gln	Ser	Glu	Glu	Leu	Lys	Glu	Gly	Trp	Asn	Lys	Thr	Glu	Thr	Ser	
225					230					235					240	
cag	gtt	gct	ccg	gca	taa	tga	SEQ ID NO: 30								741	
Gln	Val	Ala	Pro	Ala	SEQ ID NO: 31											
				245												

Figure 11B

atg Met 1	ccg Pro	cgt Arg	ggc Gly	tgg Trp 5	gcc Ala	gcc Ala	ccc Pro	ttg Leu	ctc Leu 10	ctg Leu	ctg Leu	ctg Leu	ctc Leu	cag Gln 15	gga Gly	48
ggc Gly	tgg Trp	ggc Gly	tgc Cys 20	ccc Pro	gac Asp	ctc Leu	gtc Val	tgc Cys 25	tac Tyr	acc Thr	gat Asp	tac Tyr	ctc Leu 30	cag Gln	acg Thr	96
gtc Val	atc Ile	tgc Cys 35	atc Ile	ctg Leu	gaa Glu	atg Met	tgg Trp 40	aac Asn	ctc Leu	cac His	ccc Pro	agc Ser 45	acg Thr	ctc Leu	acc Thr	144
ctt Leu	acc Thr 50	tgg Trp	caa Gln	gac Asp	cag Gln	tat Tyr 55	gaa Glu	gag Glu	ctg Leu	aag Lys	gac Asp 60	gag Glu	gcc Ala	acc Thr	tcc Ser	192
tgc Cys 65	agc Ser	ctc Leu	cac His	agg Arg	tcg Ser 70	gcc Ala	cac His	aat Asn	gcc Ala	acg Thr 75	cat His	gcc Ala	acc Thr	tac Tyr	acc Thr 80	240
tgc Cys	cac His	atg Met	gat Asp	gta Val 85	ttc Phe	cac His	ttc Phe	atg Met	gcc Ala 90	gac Asp	gac Asp	att Ile	ttc Phe	agt Ser 95	gtc Val	288
aac Asn	atc Ile	aca Thr	gac Asp 100	cag Gln	tct Ser	ggc Gly	aac Asn	tac Tyr 105	tcc Ser	cag Gln	gag Glu	tgt Cys	ggc Gly 110	agc Ser	ttt Phe	336
ctc Leu	ctg Leu	gct Ala 115	gag Glu	agc Ser	atc Ile	aag Lys	ccg Pro 120	gct Ala	ccc Pro	cct Pro	ttc Phe	aac Asn 125	gtg Val	act Thr	gtg Val	384
acc Thr	ttc Phe 130	tca Ser	gga Gly	cag Gln	tat Tyr	aat Asn 135	atc Ile	tcc Ser	tgg Trp	cgc Arg	tca Ser 140	gat Asp	tac Tyr	gaa Glu	gac Asp	432
cct Pro 145	gcc Ala	ttc Phe	tac Tyr	atg Met	ctg Leu 150	aag Lys	ggc Gly	aag Lys	ctt Leu	cag Gln 155	tat Tyr	gag Glu	ctg Leu	cag Gln	tac Tyr 160	480
agg Arg	aac Asn	cgg Arg	gga Gly	gac Asp 165	ccc Pro	tgg Trp	gct Ala	gtg Val	agt Ser 170	ccg Pro	agg Arg	aga Arg	aag Lys	ctg Leu 175	atc Ile	528
tca Ser	gtg Val	gac Asp	tca Ser 180	aga Arg	agt Ser	gtc Val	tcc Ser	ctc Leu 185	ctc Leu	ccc Pro	ctg Leu	gag Glu	ttc Phe 190	cgc Arg	aaa Lys	576
gac Asp	tcg Ser	agc Ser 195	tat Tyr	gag Glu	ctg Leu	cag Gln	gtg Val 200	cgg Arg	gca Ala	ggg Gly	ccc Pro	atg Met 205	cct Pro	ggc Gly	tcc Ser	624
tcc Ser	tac Tyr 210	cag Gln	ggg Gly	acc Thr	tgg Trp	agt Ser 215	gaa Glu	tgg Trp	agt Ser	gac Asp	ccg Pro 220	gtc Val	atc Ile	ttt Phe	cag Gln	672

Figure 12A

acc Thr 225	cag Gln	tca Ser	gag Glu	gag Glu	tta Leu 230	aag Lys	gaa Glu	ggc Gly	tgg Trp	aac Asn 235	gat Asp	gac Asp	gat Asp	gac Asp	aag Lys 240	720
ggc Gly	tcc Ser	ggc Gly	gac Asp	aaa Lys 245	act Thr	cac His	aca Thr	tgc Cys	cca Pro 250	ccg Pro	tgc Cys	cca Pro	gca Ala	cct Pro 255	gaa Glu	768
gcc Ala	ctg Leu	ggg Gly	gca Ala 260	ccg Pro	tca Ser	gtc Val	ttc Phe	ctc Leu 265	ttc Phe	ccc Pro	cca Pro	aaa Lys	ccc Pro 270	aag Lys	gac Asp	816
acc Thr	ctc Leu	atg Met 275	atc Ile	tcc Ser	cgg Arg	acc Thr	cct Pro 280	gag Glu	gtc Val	aca Thr	tgc Cys	gtg Val 285	gtg Val	gtg Val	gac Asp	864
gtg Val	agc Ser 290	cac His	gaa Glu	gac Asp	cct Pro	gag Glu 295	gtc Val	aag Lys	ttc Phe	aac Asn	tgg Trp 300	tac Tyr	gtg Val	gac Asp	ggc Gly	912
gtg Val 305	gag Glu	gtg Val	cat His	aat Asn	gcc Ala 310	aag Lys	aca Thr	aag Lys	ccg Pro	cgg Arg 315	gag Glu	gag Glu	cag Gln	tac Tyr	aac Asn 320	960
agc Ser	acg Thr	tac Tyr	cgt Arg	gtg Val 325	gtc Val	agc Ser	gtc Val	ctc Leu	acc Thr 330	gtc Val	ctg Leu	cac His	cag Gln	gac Asp 335	tgg Trp	1008
ctg Leu	aat Asn	ggc Gly	aag Lys 340	gag Glu	tac Tyr	aag Lys	tgc Cys	aag Lys 345	gtc Val	tcc Ser	aac Asn	aaa Lys	gcc Ala 350	ctc Leu	cca Pro	1056
gcc Ala	ccc Pro	atc Ile 355	gag Glu	aaa Lys	acc Thr	atc Ile	tcc Ser 360	aaa Lys	gcc Ala	aaa Lys	ggg Gly	cag Gln 365	ccc Pro	cga Arg	gaa Glu	1104
cca Pro	cag Gln 370	gtg Val	tac Tyr	acc Thr	ctg Leu	ccc Pro 375	cca Pro	tcc Ser	cgg Arg	gag Glu	gag Glu 380	atg Met	acc Thr	aag Lys	aac Asn	1152
cag Gln 385	gtc Val	agc Ser	ctg Leu	acc Thr	tgc Cys 390	ctg Leu	gtc Val	aaa Lys	ggc Gly	ttc Phe 395	tat Tyr	ccc Pro	agc Ser	gac Asp	atc Ile 400	1200
gcc Ala	gtg Val	gag Glu	tgg Trp	gag Glu 405	agc Ser	aat Asn	ggg Gly	cag Gln	ccg Pro 410	gag Glu	aac Asn	aac Asn	tac Tyr	aag Lys 415	acc Thr	1248
acg Thr	cct Pro	ccc Pro	gtg Val 420	ctg Leu	gac Asp	tcc Ser	gac Asp	ggc Gly 425	tcc Ser	ttc Phe	ttc Phe	ctc Leu	tat Tyr 430	agc Ser	aag Lys	1296
ctc Leu	acc Thr	gtg Val 435	gac Asp	aag Lys	agc Ser	agg Arg	tgg Trp 440	cag Gln	cag Gln	ggg Gly	aac Asn	gtc Val 445	ttc Phe	tca Ser	tgc Cys	1344

Figure 12B

tcc	gtg	atg	cat	gag	gct	ctg	cac	aac	cac	tac	acg	cag	aag	agc	ctc	1392
Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr	Thr	Gln	Lys	Ser	Leu	
	450					455					460					
tcc	ctg	tcc	ccg	ggt	aaa	tga	SEQ ID NO: 32									1413
Ser	Leu	Ser	Pro	Gly	Lys		SEQ ID NO: 33									
465					470											

Figure 12C

Q ID NO:34	atg	ccc	cgg	ggc	cca	gtg	gct	gcc	tta	ctc	ctg	ctg	att	ctc	cat	gga	48
Q ID NO:35	Met	Pro	Arg	Gly	Pro	Val	Ala	Ala	Leu	Leu	Leu	Leu	Ile	Leu	His	Gly	
	1				5				10					15			
	gct	tgg	agc	tgc	ctg	gac	ctc	act	tgc	tac	act	gac	tac	ctc	tgg	acc	96
	Ala	Trp	Ser	Cys	Leu	Asp	Leu	Thr	Cys	Tyr	Thr	Asp	Tyr	Leu	Trp	Thr	
				20					25					30			
	atc	acc	tgt	gtc	ctg	gag	aca	cgg	agc	ccc	aac	ccc	agc	ata	ctc	agt	144
	Ile	Thr	Cys	Val	Leu	Glu	Thr	Arg	Ser	Pro	Asn	Pro	Ser	Ile	Leu	Ser	
			35					40					45				
	ctc	acc	tgg	caa	gat	gaa	tat	gag	gaa	ctt	cag	gac	caa	gag	acc	ttc	192
	Leu	Thr	Trp	Gln	Asp	Glu	Tyr	Glu	Glu	Leu	Gln	Asp	Gln	Glu	Thr	Phe	
			50				55					60					
	tgc	agc	cta	cac	agg	tct	ggc	cac	aac	acc	aca	cat	ata	tgg	tac	acg	240
	Cys	Ser	Leu	His	Arg	Ser	Gly	His	Asn	Thr	Thr	His	Ile	Trp	Tyr	Thr	
	65					70					75					80	
	tgc	cat	atg	cgc	ttg	tct	caa	ttc	ctg	tcc	gat	gaa	gtt	ttc	att	gtc	288
	Cys	His	Met	Arg	Leu	Ser	Gln	Phe	Leu	Ser	Asp	Glu	Val	Phe	Ile	Val	
					85					90					95		
	aat	gtg	acg	gac	cag	tct	ggc	aac	aac	tcc	caa	gag	tgt	ggc	agc	ttt	336
	Asn	Val	Thr	Asp	Gln	Ser	Gly	Asn	Asn	Ser	Gln	Glu	Cys	Gly	Ser	Phe	
				100				105						110			
	gtc	ctg	gct	gag	agc	atc	aaa	cca	gct	ccc	ccc	ttg	aac	gtg	act	gtg	384
	Val	Leu	Ala	Glu	Ser	Ile	Lys	Pro	Ala	Pro	Pro	Leu	Asn	Val	Thr	Val	
			115					120					125				
	gcc	ttc	tca	gga	cgc	tat	gat	atc	tcc	tgg	gac	tca	gct	tat	gac	gaa	432
	Ala	Phe	Ser	Gly	Arg	Tyr	Asp	Ile	Ser	Trp	Asp	Ser	Ala	Tyr	Asp	Glu	
			130				135					140					
	ccc	tcc	aac	tac	gtg	ctg	agg	ggc	aag	cta	caa	tat	gag	ctg	cag	tat	480
	Pro	Ser	Asn	Tyr	Val	Leu	Arg	Gly	Lys	Leu	Gln	Tyr	Glu	Leu	Gln	Tyr	
	145					150					155					160	
	cgg	aac	ctc	aga	gac	ccc	tat	gct	gtg	agg	ccg	gtg	acc	aag	ctg	atc	528
	Arg	Asn	Leu	Arg	Asp	Pro	Tyr	Ala	Val	Arg	Pro	Val	Thr	Lys	Leu	Ile	
					165					170					175		
	tca	gtg	gac	tca	aga	aac	gtc	tct	ctt	ctc	cct	gaa	gag	ttc	cac	aaa	576
	Ser	Val	Asp	Ser	Arg	Asn	Val	Ser	Leu	Leu	Pro	Glu	Glu	Phe	His	Lys	
				180					185					190			
	gat	tct	agc	tac	cag	ctg	cag	gtg	cgg	gca	gcg	cct	cag	cca	ggc	act	624
	Asp	Ser	Ser	Tyr	Gln	Leu	Gln	Val	Arg	Ala	Ala	Pro	Gln	Pro	Gly	Thr	
			195					200					205				
	tca	ttc	agg	ggg	acc	tgg	agt	gag	tgg	agt	gac	ccc	gtc	atc	ttt	cag	672
	Ser	Phe	Arg	Gly	Thr	Trp	Ser	Glu	Trp	Ser	Asp	Pro	Val	Ile	Phe	Gln	
							215					220					

Figure 13A

acc	cag	gct	ggg	gag	ccc	gag	gca	ggc	tgg	gac	ggc	tcc	ggc	tct	aga		720
Thr	Gln	Ala	Gly	Glu	Pro	Glu	Ala	Gly	Trp	Asp	Gly	Ser	Gly	Ser	Arg		
225					230					235					240		
<hr/>																	
gagccccgcg gaccgacaat caagccctgt cctccatgca aatgcccgagg taagtacta																780	
gaccagagct ccactcccgg gagaatggta agtgctataa acatccctgc actagaggat																840	
aagccatgta cagatccatt tccatctctc ctcatcagca cctaacctcg aggggtggacc																900	
atccgtcttc atcttccctc caaagatcaa ggatgtactc atgatctccc tgagcccat																960	
agtcacatgt gtgggtgggtg atgtgagcga ggatgacca gatgtccaga tcagctggtt																1020	
tgtgaacaac gtggaagtac acacagctca gacacaaacc catagagagg attacaacag																1080	
tactctccgg gtgggtcagtg ccctcccat ccagcaccag gactggatga gtggcaaggc																1140	
tttcgcatgc gccgtcaaca acaaagacct ccagcgccc atcgagagaa ccatctcaaa																1200	
acccaaaggt gagagctgca gcctgactgc atgggggctg ggatgggcat aaggataaag																1260	
gtctgtgtgg acagccttct gcttcagcca tgacctttgt gtatgtttct accctcacag																1320	
ggtcagtaaag agctccacag gtatatgtct tgccctccacc agaagaagag atgactaaga																1380	
aacaggtcac tctgacctgc atgggtcacag acttcatgcc tgaagacatt tacgtggagt																1440	
ggaccaaaaa cgaggaaaaca gagctaaact acaagaacac tgaaccagtc ctggactctg																1500	
atggttctta cttcatgtac agcaagctga gagtggaaaa gaagaactgg gtggaaagaa																1560	
atagctactc ctgttcagtg gtccacgagg gtctgcacaa tcaccacacg actaagagct																1620	
tctcccgac tccgggtaaa tgagctcagc acccacaanaa ctctcaggtc caaagagaca																1680	
cccacactca tctccatgct tcccttgat aaataaagca cccagcaatg cctgggacca																1740	
tgtaatagga attc																1754	

Figure 13B

ctgcaggtcg	acaccacc	atg	ccc	cgg	ggc	cca	gtg	gct	gcc	tta	ctc	ctg	51
		Met	Pro	Arg	Gly	Pro	Val	Ala	Ala	Leu	Leu	Leu	
		1				5				10			
ctg	att	ctc	cat	gga	gct	tgg	agc	tgc	ctg	gac	ctc	act	99
Leu	Ile	Leu	His	Gly	Ala	Trp	Ser	Cys	Leu	Asp	Leu	Thr	
			15					20				25	
gac	tac	ctc	tgg	acc	atc	acc	tgt	gtc	ctg	gag	aca	cgg	147
Asp	Tyr	Leu	Trp	Thr	Ile	Thr	Cys	Val	Leu	Glu	Thr	Arg	
		30					35					40	
ccc	agc	ata	ctc	agt	ctc	acc	tgg	caa	gat	gaa	tat	gag	195
Pro	Ser	Ile	Leu	Ser	Leu	Thr	Trp	Gln	Asp	Glu	Tyr	Glu	
	45					50					55	Glu	
gac	caa	gag	acc	ttc	tgc	agc	cta	cac	agg	tct	ggc	cac	243
Asp	Gln	Glu	Thr	Phe	Cys	Ser	Leu	His	Arg	Ser	Gly	His	
60					65					70		Aasn	
cat	ata	tgg	tac	acg	tgc	cat	atg	cgc	ttg	tct	caa	ttc	291
His	Ile	Trp	Tyr	Thr	Cys	His	Met	Arg	Leu	Ser	Gln	Phe	
				80					85			Leu	
gaa	gtt	ttc	att	gtc	aat	gtg	acg	gac	cag	tct	ggc	aac	339
Glu	Val	Phe	Ile	Val	Asn	Val	Thr	Asp	Gln	Ser	Gly	Asn	
			95					100				Aasn	
gag	tgt	ggc	agc	ttt	gtc	ctg	gct	gag	agc	atc	aaa	cca	387
Glu	Cys	Gly	Ser	Phe	Val	Leu	Ala	Glu	Ser	Ile	Lys	Pro	
		110					115					Ala	
ttg	aac	gtg	act	gtg	gcc	ttc	tca	gga	cgc	tat	gat	atc	435
Leu	Asn	Val	Thr	Val	Ala	Phe	Ser	Gly	Arg	Tyr	Asp	Ile	
	125					130					135	Ser	
tca	gct	tat	gac	gaa	ccc	tcc	aac	tac	gtg	ctg	agg	ggc	483
Ser	Ala	Tyr	Asp	Glu	Pro	Ser	Asn	Tyr	Val	Leu	Arg	Gly	
140					145					150		Aag	
tat	gag	ctg	cag	tat	cgg	aac	ctc	aga	gac	ccc	tat	gct	531
Tyr	Glu	Leu	Gln	Tyr	Arg	Asn	Leu	Arg	Asp	Pro	Tyr	Ala	
				160					165			Val	
gtg	acc	aag	ctg	atc	tca	gtg	gac	tca	aga	aac	gtc	tct	579
Val	Thr	Lys	Leu	Ile	Ser	Val	Asp	Ser	Arg	Asn	Val	Ser	
			175					180				Leu	
gaa	gag	ttc	cac	aaa	gat	tct	agc	tac	cag	ctg	cag	gtg	627
Glu	Glu	Phe	His	Lys	Asp	Ser	Ser	Tyr	Gln	Leu	Gln	Val	
		190					195					Arg	
cct	cag	cca	ggc	act	tca	ttc	agg	ggg	acc	tgg	agt	gag	675
Pro	Gln	Pro	Gly	Thr	Ser	Phe	Arg	Gly	Thr	Trp	Ser	Glu	
	205					210					215	Trp	

Figure 14A

ccc	gtc	atc	ttt	cag	acc	cag	gct	ggg	gag	ccc	gag	gca	ggc	tgg	gac	723
Pro	Val	Ile	Phe	Gln	Thr	Gln	Ala	Gly	Glu	Pro	Glu	Ala	Gly	Trp	Asp	
220					225					230					235	
ggc	agc	gga	cac	cac	cat	cat	cac	cac	ggt	agc	ggc	gac	tat	aaa	gac	771
Gly	Ser	Gly	His	His	His	His	His	His	Gly	Ser	Gly	Asp	Tyr	Lys	Asp	
				240					245					250		
gat	gac	gat	aag	tagtgagaat	tc	SEQ ID NO: 36										795
Asp	Asp	Asp	Lys	SEQ ID NO: 37												
			255													

Figure 14B

atg Met 1	aaa Lys	ttc Phe	tta Leu	gtc Val 5	aac Asn	gtt Val	gcc Ala	ctt Leu	gtt Val 10	ttt Phe	atg Met	gtc Val	gtg Val	tac Tyr 15	att Ile	48
tct Ser	tac Tyr	atc Ile	tat Tyr 20	gcc Ala	ggc Gly	agc Ser	gga Gly	cac His 25	cac His	cat His	cat His	cac His	cac His 30	ggg Gly	agc Ser	96
ggc Gly	gac Asp	tat Tyr 35	aaa Lys	gac Asp	gat Asp	gac Asp	gat Asp 40	aag Lys	ggg Gly	tcc Ser	gga Gly	tgc Cys 45	ctg Leu	gac Asp	ctc Leu	144
act Thr	tgc Cys 50	tac Tyr	act Thr	gac Asp	tac Tyr	ctc Leu 55	tgg Trp	acc Thr	atc Ile	acc Thr	tgt Cys 60	gtc Val	ctg Leu	gag Glu	aca Thr	192
cgg Arg 65	agc Ser	ccc Pro	aac Asn	ccc Pro	agc Ser 70	ata Ile	ctc Leu	agt Ser	ctc Leu	acc Thr 75	tgg Trp	caa Gln	gat Asp	gaa Glu	tat Tyr 80	240
gag Glu	gaa Glu	ctt Leu	cag Gln	gac Asp 85	caa Gln	gag Glu	acc Thr	ttc Phe	tgc Cys 90	agc Ser	cta Leu	cac His	agg Arg	tct Ser 95	ggc Gly	288
cac His	aac Asn	acc Thr	aca Thr 100	cat His	ata Ile	tgg Trp	tac Tyr	acg Thr 105	tgc Cys	cat His	atg Met	cgc Arg	ttg Leu 110	tct Ser	caa Gln	336
ttc Phe	ctg Leu	tcc Ser 115	gat Asp	gaa Glu	gtt Val	ttc Phe	att Ile 120	gtc Val	aat Asn	gtg Val	acg Thr	gac Asp 125	cag Gln	tct Ser	ggc Gly	384
aac Asn 130	aac Asn	tcc Ser	caa Gln	gag Glu	tgt Cys	ggc Gly 135	agc Ser	ttt Phe	gtc Val	ctg Leu	gct Ala 140	gag Glu	agc Ser	atc Ile	aaa Lys	432
cca Pro 145	gct Ala	ccc Pro	ccc Pro	ttg Leu	aac Asn 150	gtg Val	act Thr	gtg Val	gcc Ala	ttc Phe 155	tca Ser	gga Gly	cgc Arg	tat Tyr	gat Asp 160	480
atc Ile	tcc Ser	tgg Trp	gac Asp	tca Ser 165	gct Ala	tat Tyr	gac Asp	gaa Glu	ccc Pro 170	tcc Ser	aac Asn	tac Tyr	gtg Val	ctg Leu 175	agg Arg	528
ggc Gly	aag Lys	cta Leu	caa Gln 180	tat Tyr	gag Glu	ctg Leu	cag Gln	tat Tyr 185	cgg Arg	aac Asn	ctc Leu	aga Arg	gac Asp 190	ccc Pro	tat Tyr	576
gct Ala	gtg Val	agg Arg 195	ccg Pro	gtg Val	acc Thr	aag Lys	ctg Leu 200	atc Ile	tca Ser	gtg Val	gac Asp	tca Ser 205	aga Arg	aac Asn	gtc Val	624
tct Ser	ctt Leu 210	ctc Leu	cct Pro	gaa Glu	gag Glu	ttc Phe 215	cac His	aaa Lys	gat Asp	tct Ser	agc Ser 220	tac Tyr	cag Gln	ctg Leu	cag Gln	672

Figure 15A

gtg	cgg	gca	gcg	cct	cag	cca	ggc	act	tca	ttc	agg	ggg	acc	tgg	agt	720
Val	Arg	Ala	Ala	Pro	Gln	Pro	Gly	Thr	Ser	Phe	Arg	Gly	Thr	Trp	Ser	
225					230					235					240	
gag	tgg	agt	gac	ccc	gtc	atc	ttt	cag	acc	cag	gct	ggg	gag	ccc	gag	768
Glu	Trp	Ser	Asp	Pro	Val	Ile	Phe	Gln	Thr	Gln	Ala	Gly	Glu	Pro	Glu	
				245					250					255		
gca	ggc	tgg	gac	tagtgagaat	tc	SEQ ID NO: 38										792
Ala	Gly	Trp	Asp			SEQ ID NO: 39										
			260													

Figure 15B

Timetable for the CIA Model

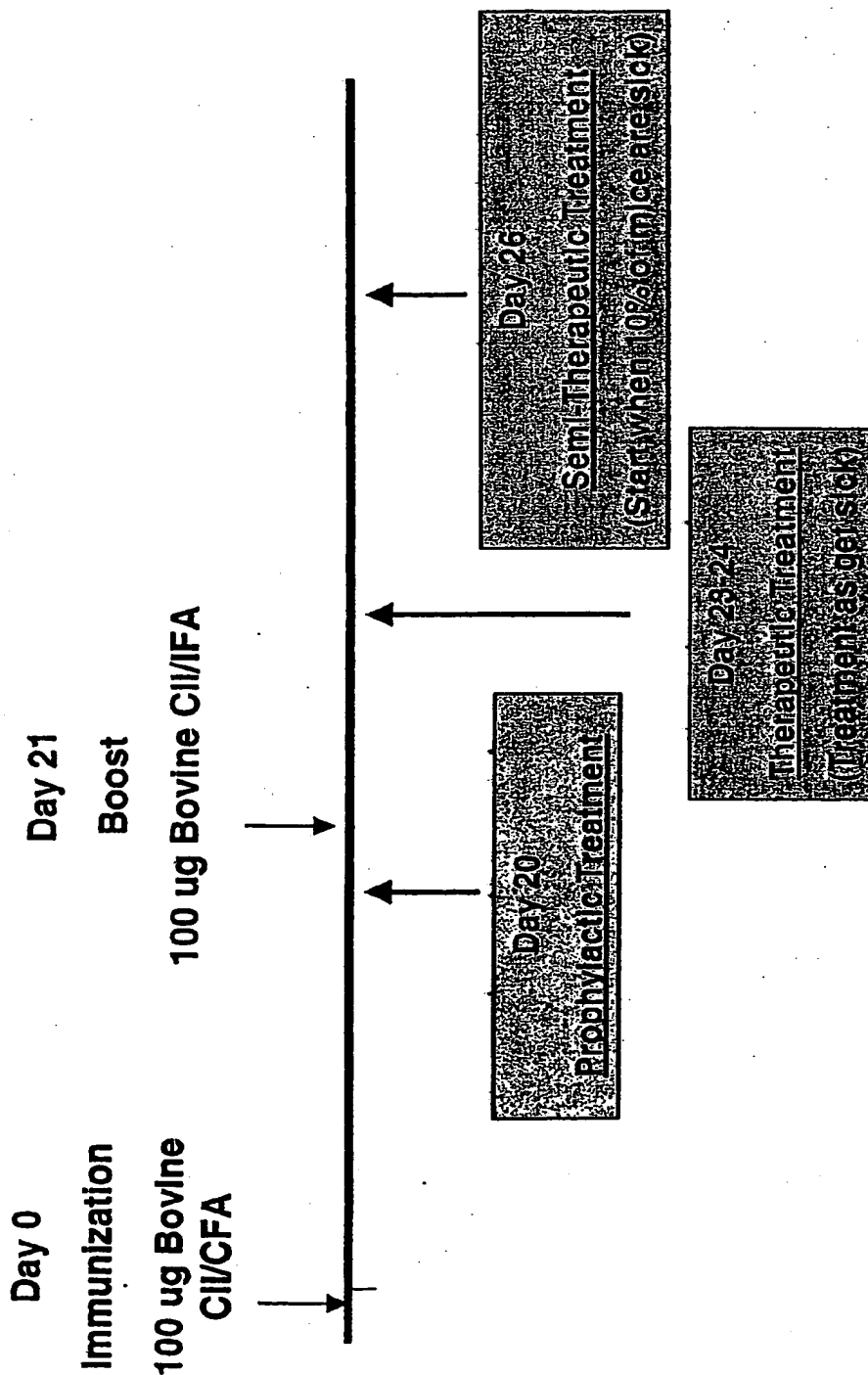


FIG.16

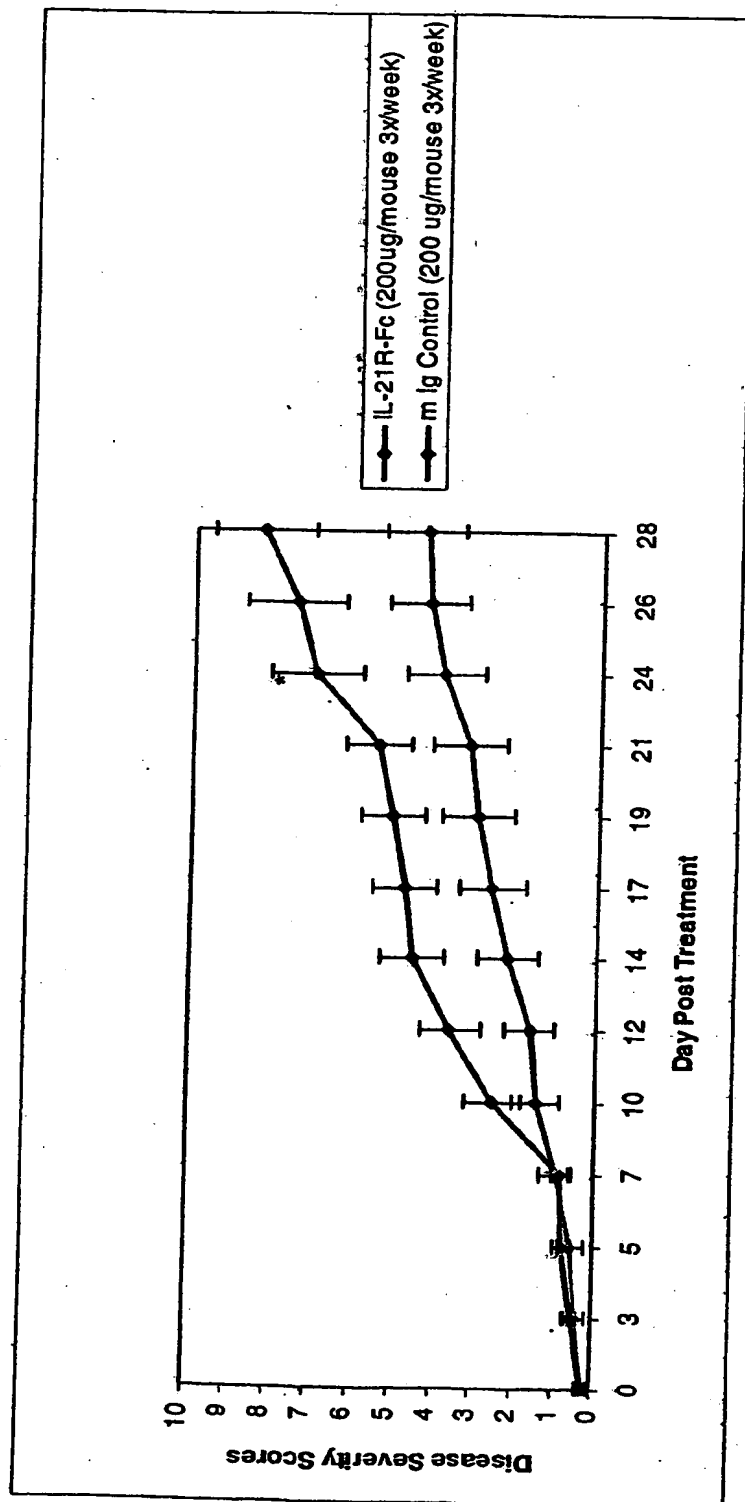


FIG.17

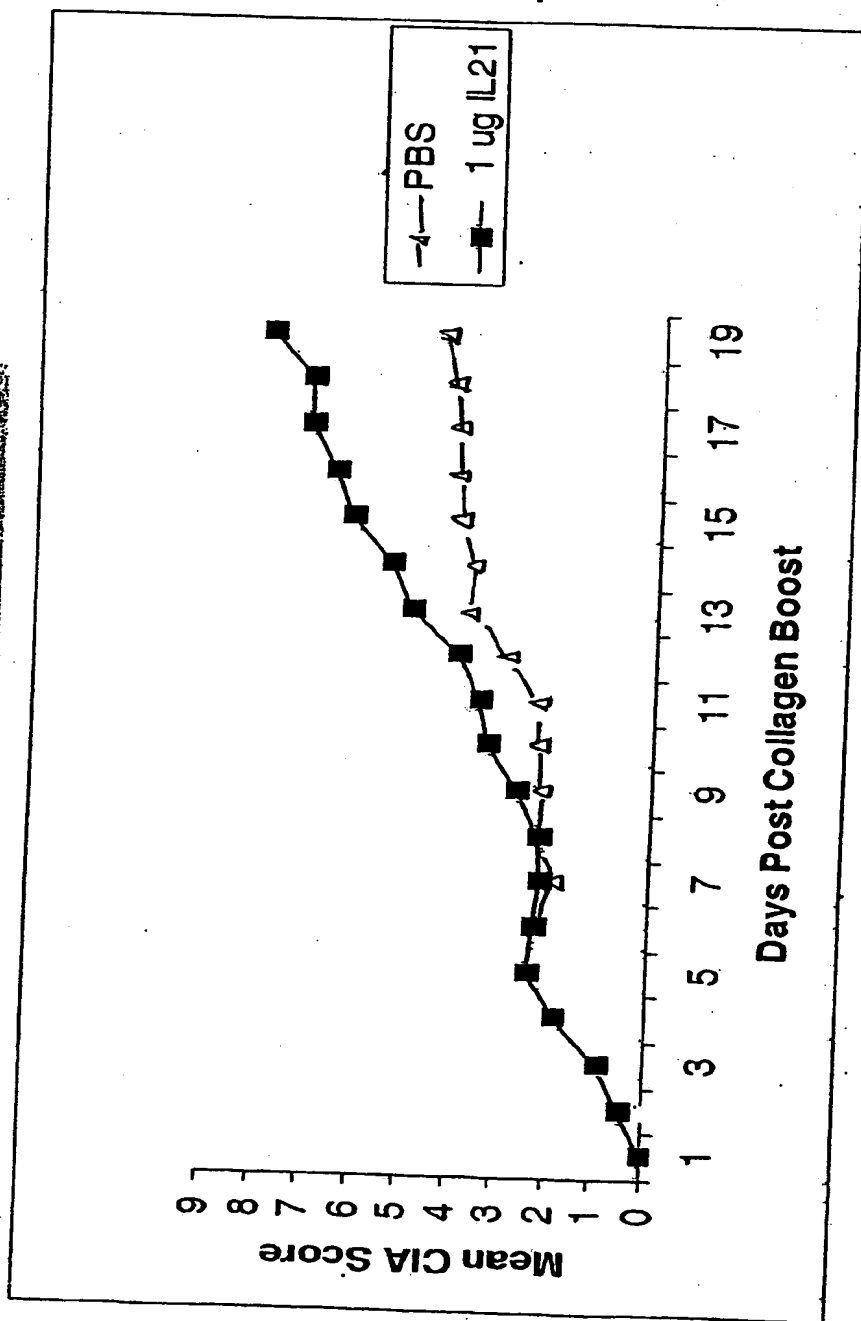


FIG.18

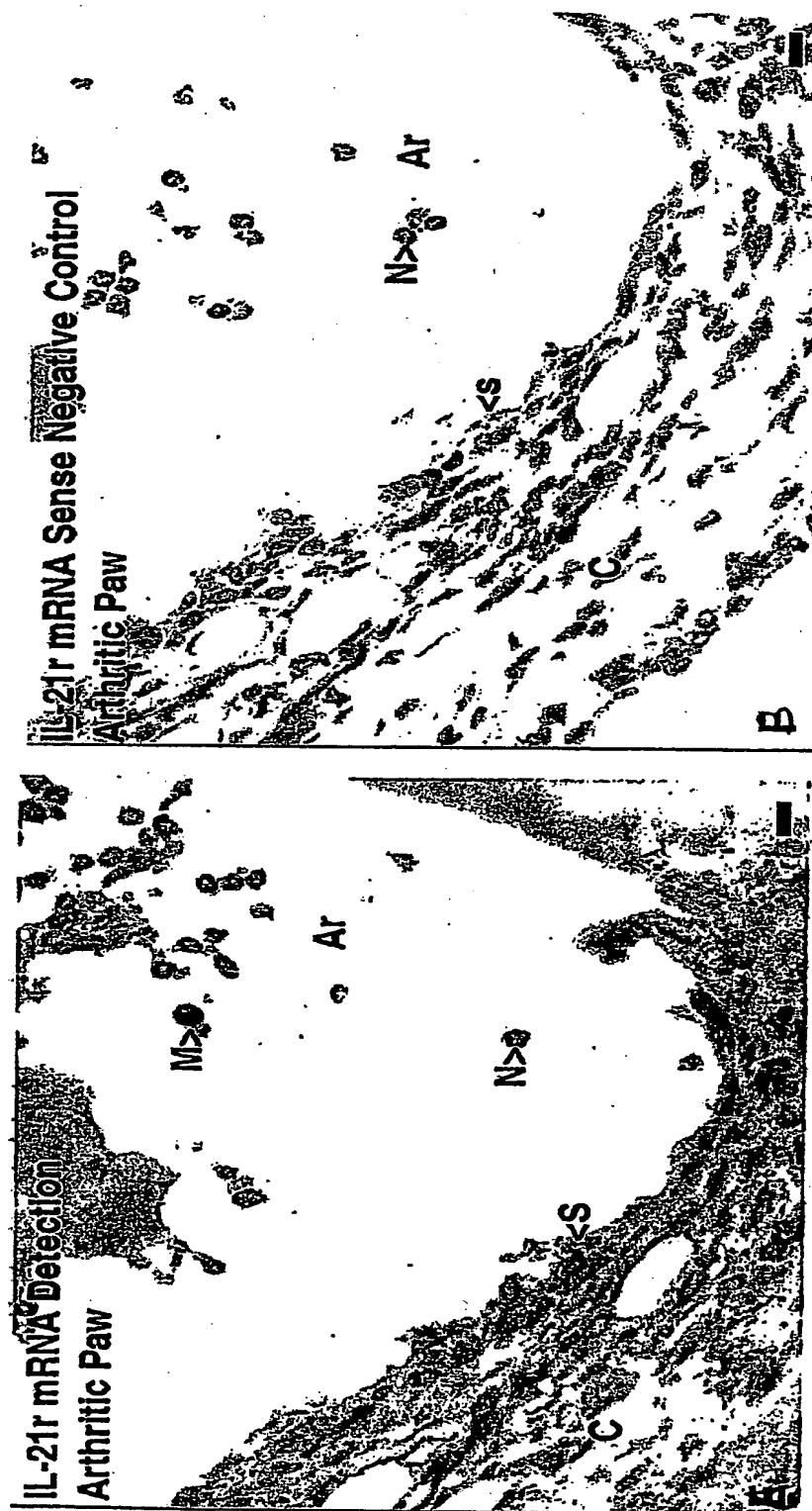
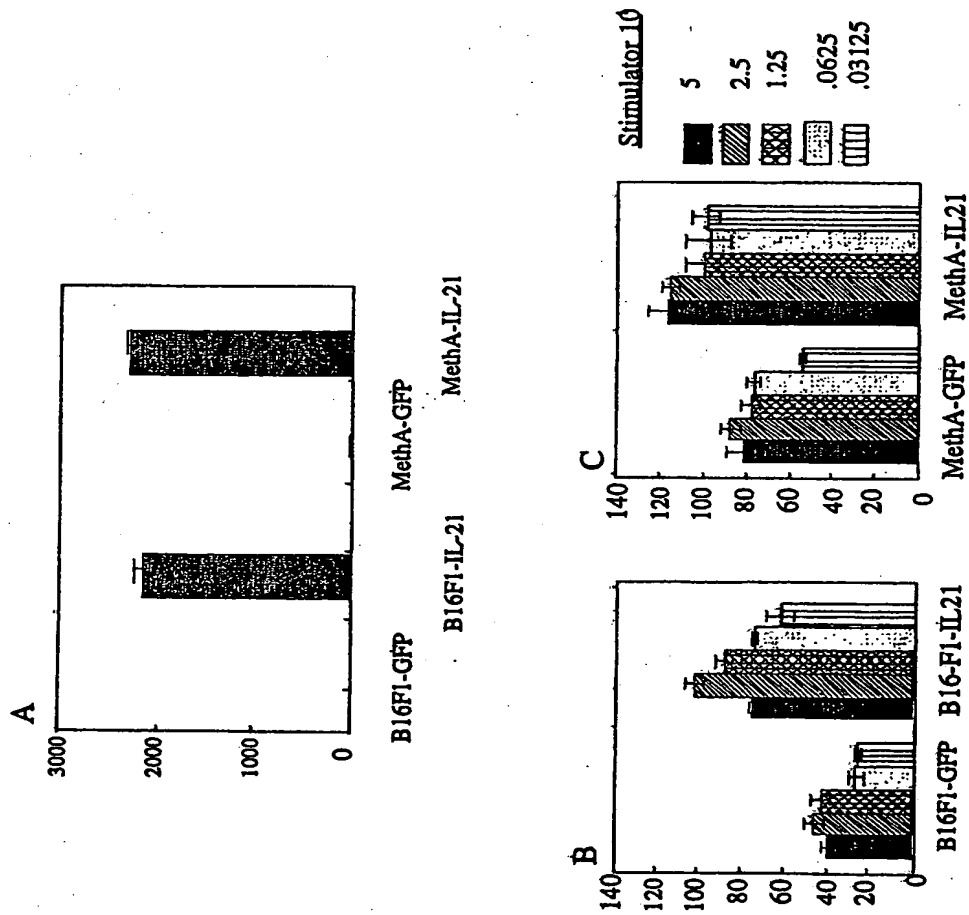
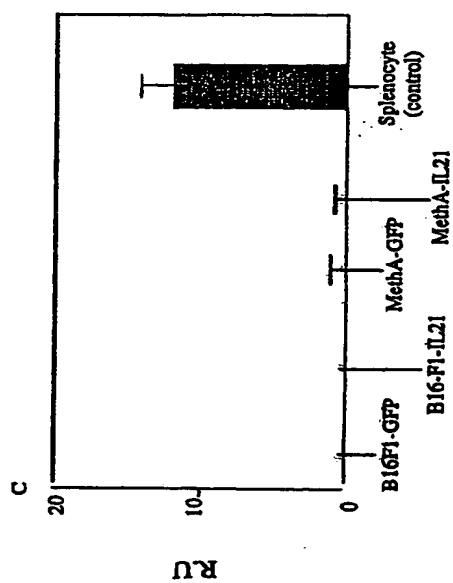


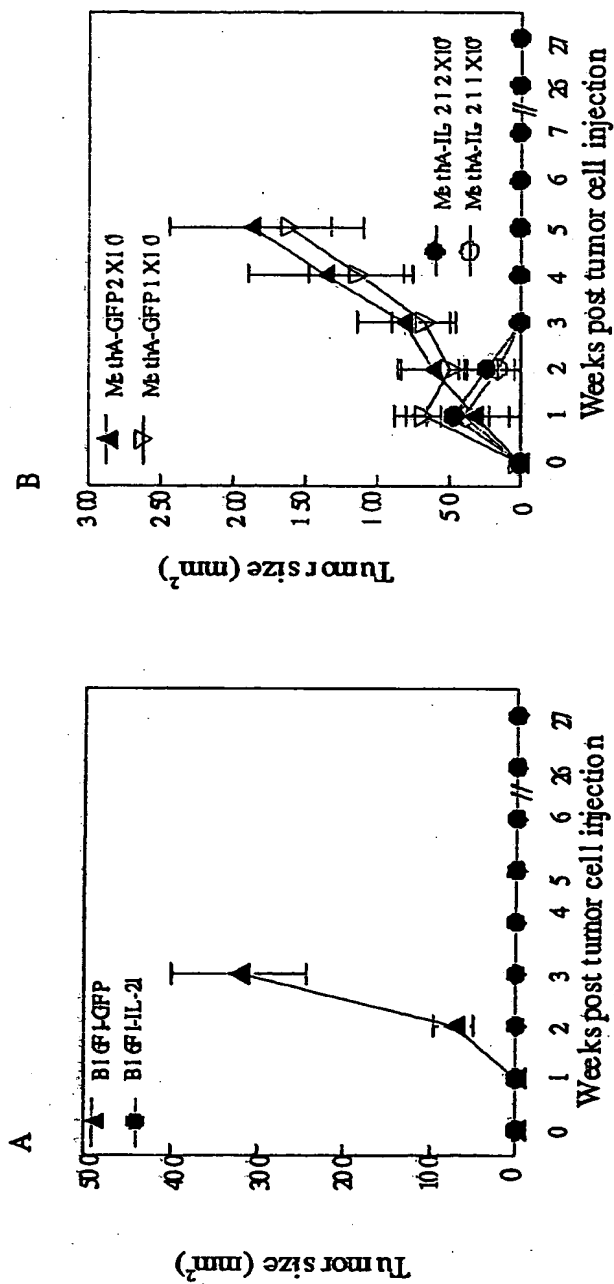
FIG.19



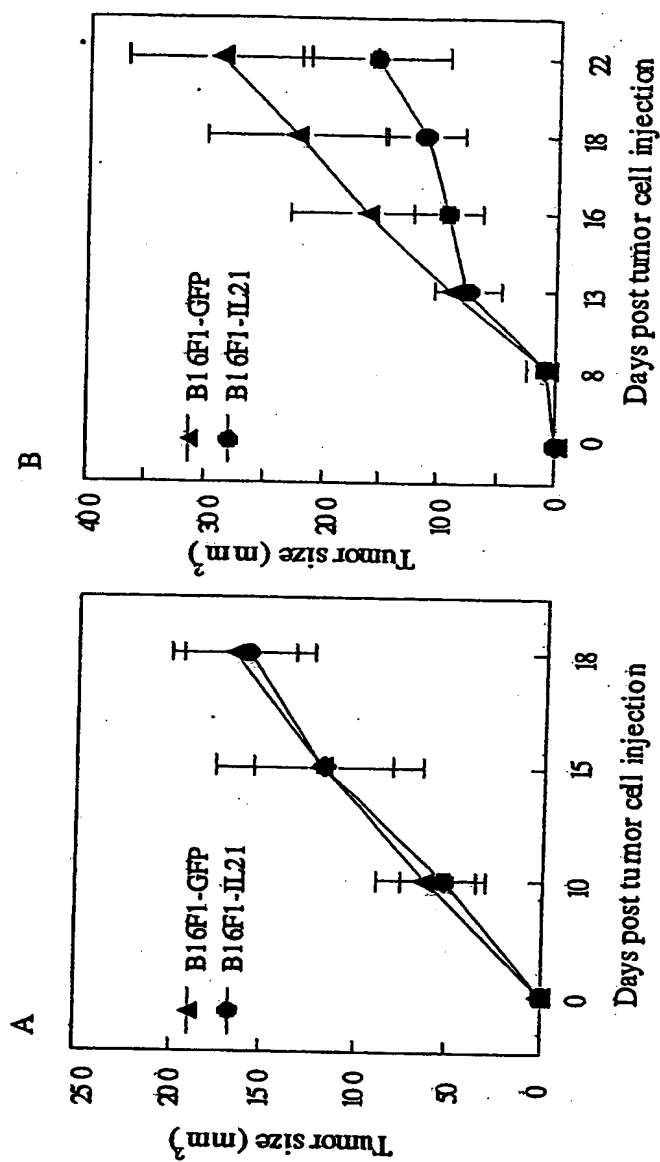
Figures 20A-20C



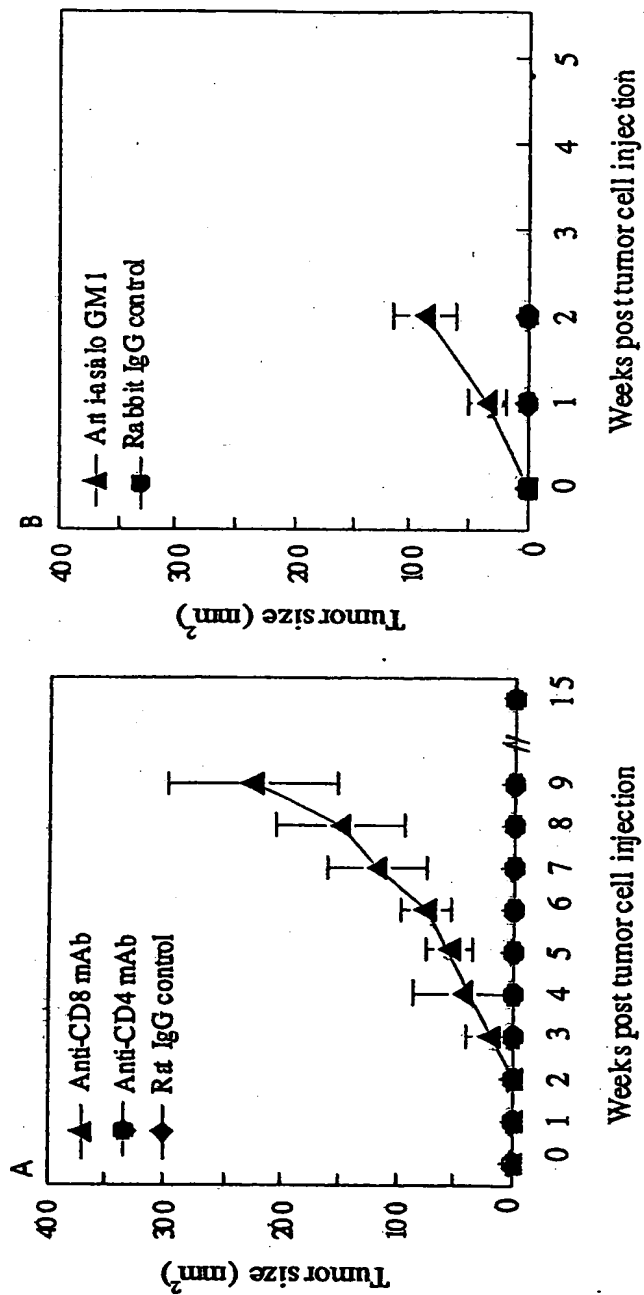
Figures 21A-21C



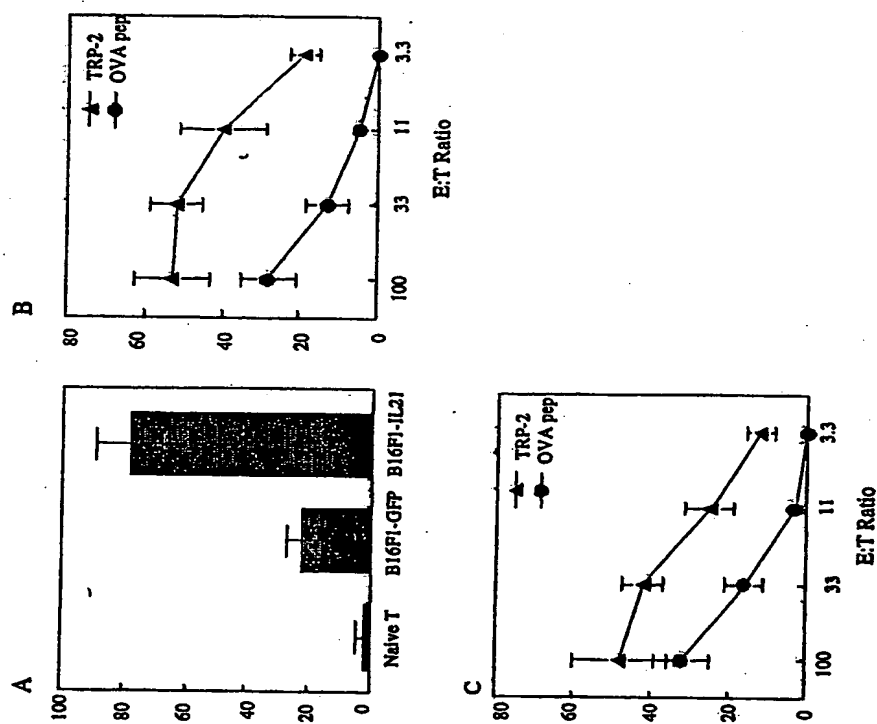
Figures 22A-22B



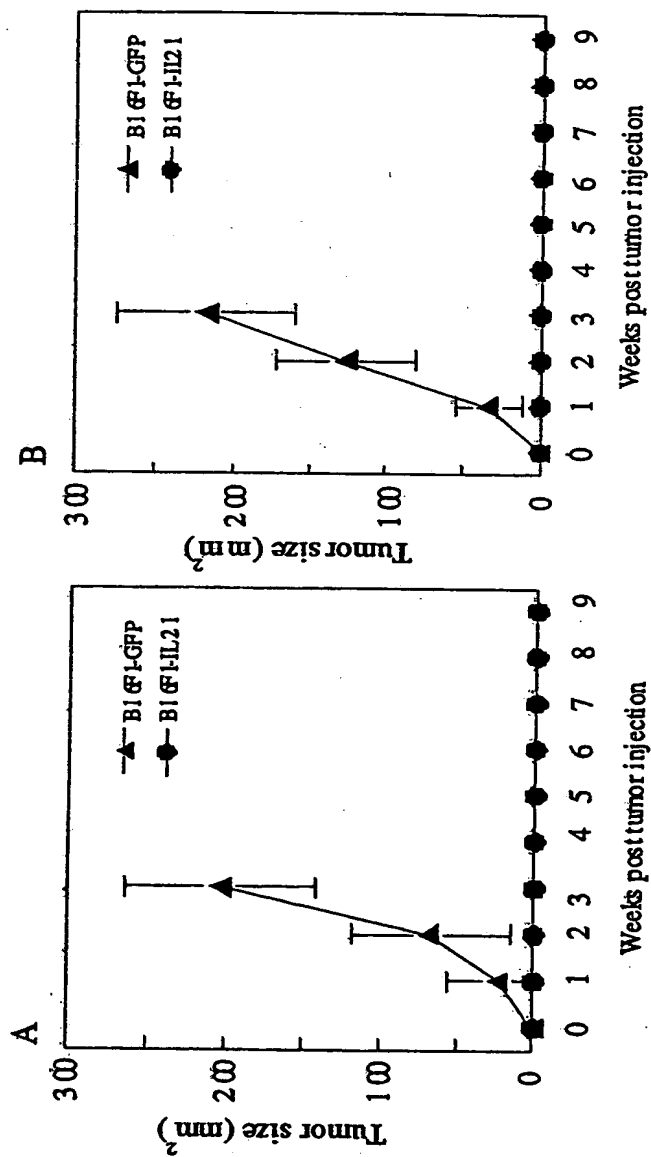
Figures 23A-23B



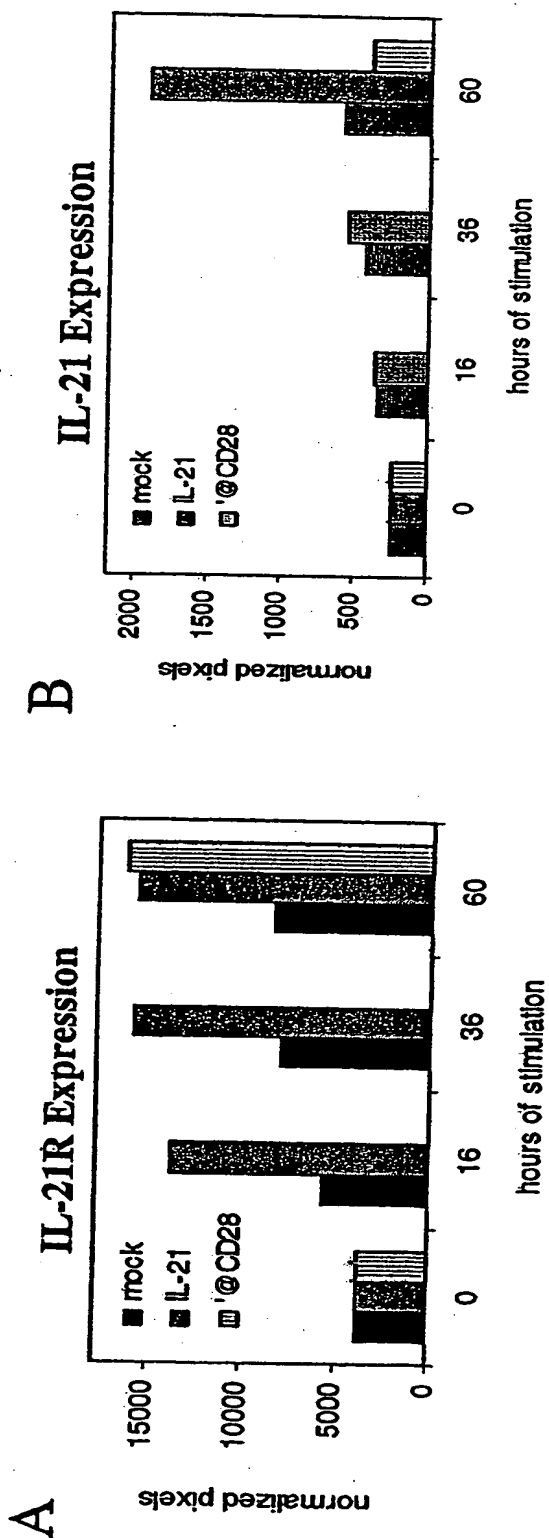
Figures 24A-24B



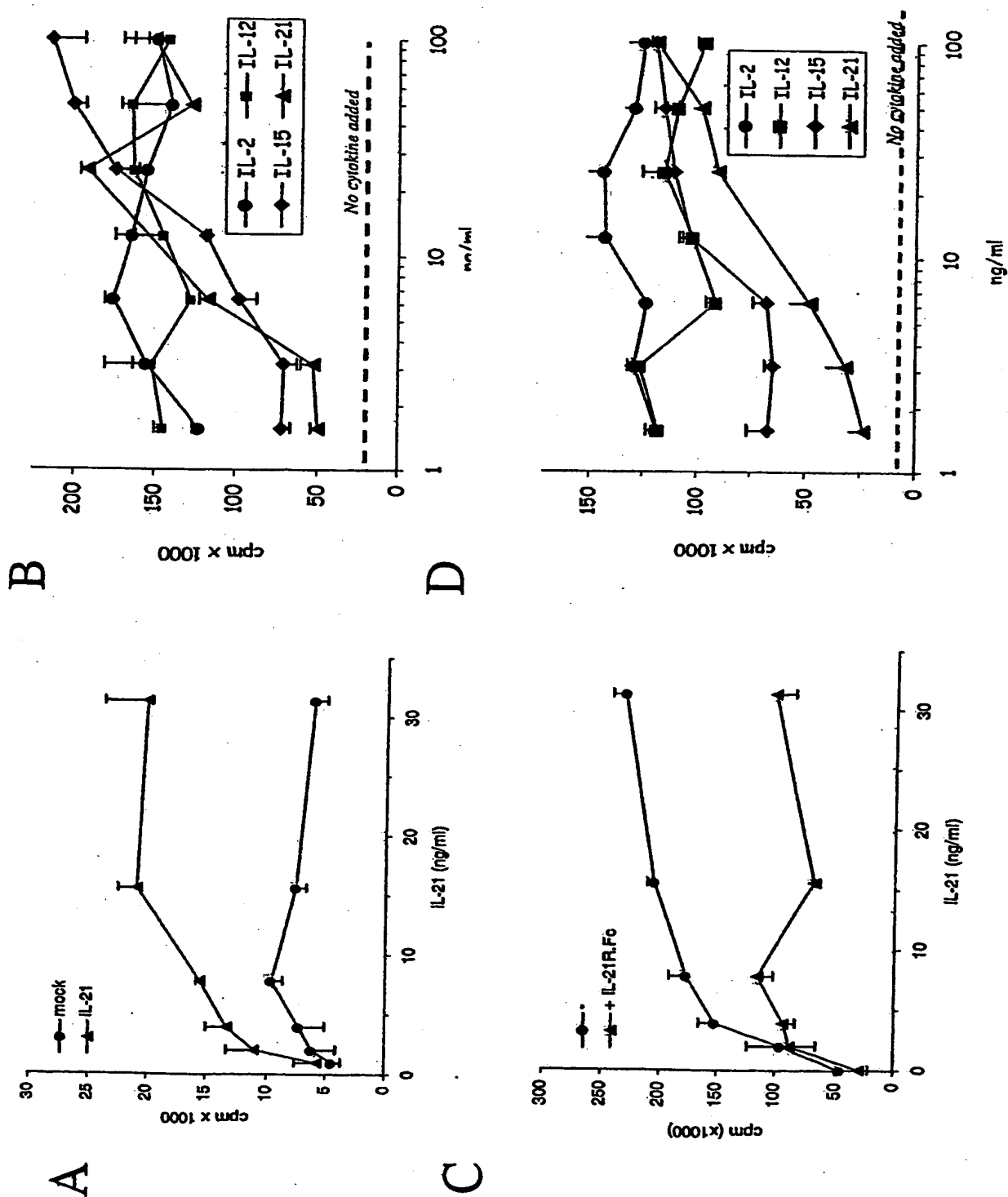
Figures 25A-25C



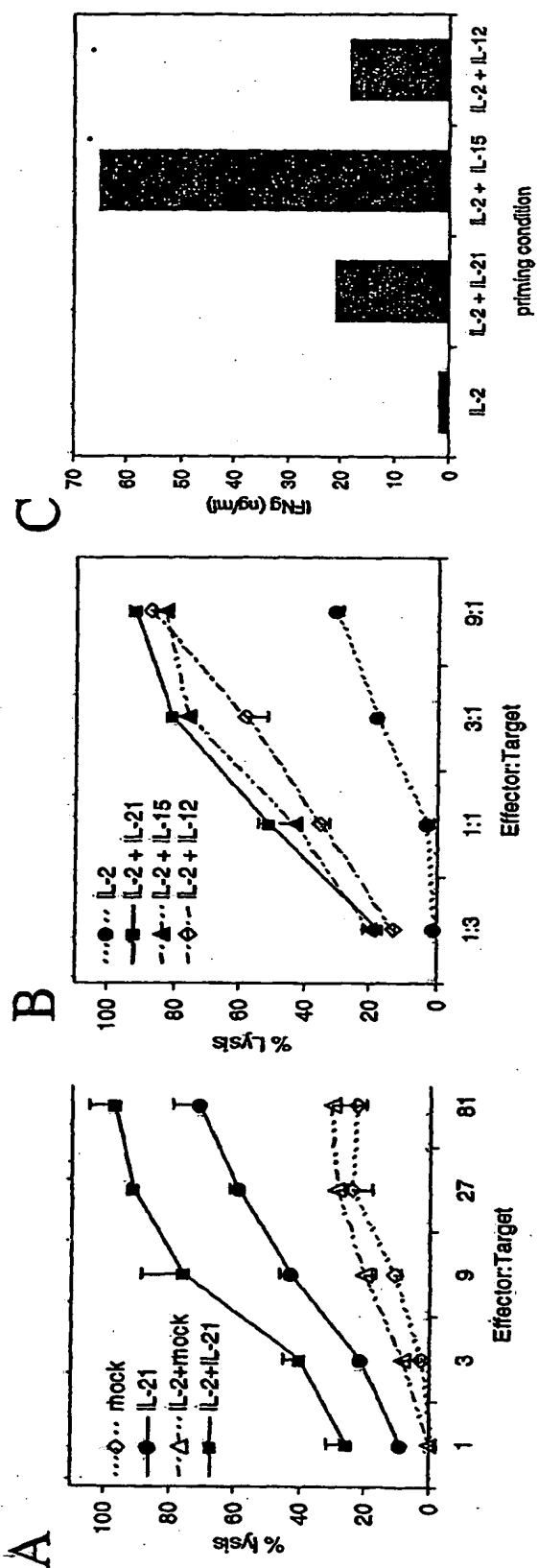
Figures 26A-26B



Figures 27A-27B



Figures 28A-28D



Figures 29A-29C

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<130> GI5320-PCT-P2

<140> Not Yet Assigned

<141> 2002-10-04

<150> 09/972,218

<151> 2001-10-04

<160> 44

<170> PatentIn version 3.1

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<211> 2665

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<400> 1

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acacggaatg gatttctgag aaagaagccg aaacagaagg cccgtgggag tcagcatgcc      240
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ctttcagacc cagtcagagg agttaaagga aggctggaac cctcacctgc tgcttctcct      960
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gggctgcagc ggagacttca agaaatgggt ggggtgaccc ttactggct ccagcctgga      1140
gctgggaccc tggagccag aggtgccctc caccctggag gtgtacagct gccacccacc      1200
acggagcccc gccaaaggc tgcagctcac ggagctacaa gaaccagcag agctgggtgga      1260
gtctgacggt gtgcccaagc ccagcttctg gccgacagcc cagaactcgg ggggctcagc      1320

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 gaccacagtc ctgtcctgtg gctgtgtctc agctggcagc cctgggctag gaggggccct 1560
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 tcaaaaaaaaa aaaaaaaaaat ctaga 2665

<210> 2
 <211> 538
 <212> PRT
 <213> Human

<400> 2

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 1 5 10 15
 Gly Trp Gly Cys Pro Asp Leu Val Cys Tyr Thr Asp Tyr Leu Gln Thr
 20 25 30
 Val Ile Cys Ile Leu Glu Met Trp Asn Leu His Pro Ser Thr Leu Thr
 35 40 45
 Leu Thr Trp Gln Asp Gln Tyr Glu Glu Leu Lys Asp Glu Ala Thr Ser
 50 55 60
 Cys Ser Leu His Arg Ser Ala His Asn Ala Thr His Ala Thr Tyr Thr
 65 70 75 80

Cys His Met Asp Val Phe His Phe Met Ala Asp Asp Ile Phe Ser Val
 85 90 95
 Asn Ile Thr Asp Gln Ser Gly Asn Tyr Ser Gln Glu Cys Gly Ser Phe
 100 105 110
 Leu Leu Ala Glu Ser Ile Lys Pro Ala Pro Pro Phe Asn Val Thr Val
 115 120 125
 Thr Phe Ser Gly Gln Tyr Asn Ile Ser Trp Arg Ser Asp Tyr Glu Asp
 130 135 140
 Pro Ala Phe Tyr Met Leu Lys Gly Lys Leu Gln Tyr Glu Leu Gln Tyr
 145 150 155 160
 Arg Asn Arg Gly Asp Pro Trp Ala Val Ser Pro Arg Arg Lys Leu Ile
 165 170 175
 Ser Val Asp Ser Arg Ser Val Ser Leu Leu Pro Leu Glu Phe Arg Lys
 180 185 190
 Asp Ser Ser Tyr Glu Leu Gln Val Arg Ala Gly Pro Met Pro Gly Ser
 195 200 205
 Ser Tyr Gln Gly Thr Trp Ser Glu Trp Ser Asp Pro Val Ile Phe Gln
 210 215 220
 Thr Gln Ser Glu Glu Leu Lys Glu Gly Trp Asn Pro His Leu Leu Leu
 225 230 235 240
 Leu Leu Leu Leu Val Ile Val Phe Ile Pro Ala Phe Trp Ser Leu Lys
 245 250 255
 Thr His Pro Leu Trp Arg Leu Trp Lys Lys Ile Trp Ala Val Pro Ser
 260 265 270
 Pro Glu Arg Phe Phe Met Pro Leu Tyr Lys Gly Cys Ser Gly Asp Phe
 275 280 285
 Lys Lys Trp Val Gly Ala Pro Phe Thr Gly Ser Ser Leu Glu Leu Gly
 290 295 300
 Pro Trp Ser Pro Glu Val Pro Ser Thr Leu Glu Val Tyr Ser Cys His
 305 310 315 320
 Pro Pro Arg Ser Pro Ala Lys Arg Leu Gln Leu Thr Glu Leu Gln Glu
 325 330 335
 Pro Ala Glu Leu Val Glu Ser Asp Gly Val Pro Lys Pro Ser Phe Trp
 340 345 350
 Pro Thr Ala Gln Asn Ser Gly Gly Ser Ala Tyr Ser Glu Glu Arg Asp
 355 360 365
 Arg Pro Tyr Gly Leu Val Ser Ile Asp Thr Val Thr Val Leu Asp Ala
 370 375 380
 Glu Gly Pro Cys Thr Trp Pro Cys Ser Cys Glu Asp Asp Gly Tyr Pro
 385 390 395 400
 Ala Leu Asp Leu Asp Ala Gly Leu Glu Pro Ser Pro Gly Leu Glu Asp
 405 410 415
 Pro Leu Leu Asp Ala Gly Thr Thr Val Leu Ser Cys Gly Cys Val Ser
 420 425 430
 Ala Gly Ser Pro Gly Leu Gly Gly Pro Leu Gly Ser Leu Leu Asp Arg
 435 440 445

Leu Lys Pro Pro Leu Ala Asp Gly Glu Asp Trp Ala Gly Gly Leu Pro
 450 455 460
 Trp Gly Gly Arg Ser Pro Gly Gly Val Ser Glu Ser Glu Ala Gly Ser
 465 470 475 480
 Pro Leu Ala Gly Leu Asp Met Asp Thr Phe Asp Ser Gly Phe Val Gly
 485 490 495
 Ser Asp Cys Ser Ser Pro Val Glu Cys Asp Phe Thr Ser Pro Gly Asp
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 Glu Gly Pro Pro Arg Ser Tyr Leu Arg Gln Trp Val Val Ile Pro Pro
 515 520 525
 Pro Leu Ser Ser Pro Gly Pro Gln Ala Ser
 530 535

<210> 3
 <211> 70
 <212> PRT
 <213> Human

<400> 3

Leu Met Thr Asn Ala Phe Ile Ser Ile Ile Asp Asp Leu Ser Lys Tyr
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 20 25 30
 Leu Trp Ser Glu Trp Ser Gln Pro Ile Tyr Val Gly Asn Asp Glu His
 35 40 45
 Lys Pro Leu Arg Glu Trp Phe Val Ile Val Ile Met Ala Thr Ile Cys
 50 55 60
 Phe Ile Leu Leu Ile Leu
 65 70

<210> 4
 <211> 25
 <212> DNA
 <213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: PCR Primer

<400> 4

gagtcgagg agaaagctga tctca

25

<210> 5
 <211> 24
 <212> DNA
 <213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: PCR Primer

<400> 5

gaaagatgac cgggtcactc catt

24

<210> 6
 <211> 29
 <212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: Labelled Hybridization Oligonucleotide

<400> 6

actcgagcta tgagctgcag gtgcgggca

29

<210> 7

<211> 29

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: NN14-1b (MU-1) Labelled Hybridization Oligonucleotide

<400> 7

actcgagcta tgagctgcag gtgcgggca

29

<210> 8

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: Motif Characteristic of the Hematopoietin Receptor Family

<220>

<221> MISC_FEATURE

<222> (3)..(3)

<223> wherein Xaa is any amino acid.

<400> 8

Trp Ser Xaa Trp Ser
1 5

<210> 9

<211> 2628

<212> DNA

<213> Mouse

<400> 9

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gcttgtctga ccgcaagtct gcccatccct ggggcagcca actggcctca gcccgtgccc	120
caggcgtgcc ctgtctctgt ctggctgccc cagccctact gtcttcctct gtgtaggctc	180
tgcccagatg cccggctggt cctcagcctc aggactatct cagcagtgaac tcccctgatt	240
ctggacttgc acctgactga actcctgccc acctcaaacc ttcacctccc accaccacca	300
ctccgagtcc cgctgtgact tccacgccc ggagaccacc caagtgtccc agcctaaaga	360
atggctttct gagaaagacc ctgaaggagt aggtctggga cacagcatgc cccggggccc	420
actggctgcc ttactcctgc tgattctcca tggagcttgg agctgcctgg acctcacttg	480
ctacactgac tacctctgga ccatcacctg tgtcctggag acacggagcc ccaagcccag	540
catactcagt ctacactggc aagatgaata tgaggaactt caggaccaag agaecttctg	600

cagcctacac aggtctggcc acaacaccac acatatatgg tacacgtgcc atatgcgctt	660
gtctcaattc ctgtccgatg aagttttcat tgtcaatgtg acggaccagt ctggcaacaa	720
ctccaagag tgtggcagct ttgtcctggc tgagagcatc aaaccagctc ccccttgaa	780
cgtgactgtg gccttctcag gacgctatga tatctcctgg gactcagctt atgacgaacc	840
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gatatgggca ccagtgccca cccctgagag tttcttcag cccctgtaca gggagcacag	1260
cgggaacttc aagaaatggg ttaatacccc ttacaggcc tccagcatag agttggtgcc	1320
acagagttcc acaacaacat cagccttaca tctgtcattg tatccagcca aggagaagaa	1380
gttccccggg ctgccgggtc tggaagagca actggagtgt gatggaatgt ctgagcctgg	1440
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agaccggcca tatggtctgg tgtccattga cacagtgact gtgggagatg cagagggcct	1560
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tgggtcccca ggagggggct ctgagagtga agcagggttc cccctggtc tggacatgga	1860
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aggggagagg gaccatgagc ctgtggccag gagaaacagc aagtatctgg ggtacactgg	2520
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caaagccaat taaaataaca aaaaaaaaaa aaaagcggcc gctctaga	2628

<210> 10
 <211> 529
 <212> PRT
 <213> Mouse

<400> 10

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 20 25 30
 Ile Thr Cys Val Leu Glu Thr Arg Ser Pro Asn Pro Ser Ile Leu Ser
 35 40 45
 Leu Thr Trp Gln Asp Glu Tyr Glu Glu Leu Gln Asp Gln Glu Thr Phe
 50 55 60
 Cys Ser Leu His Arg Ser Gly His Asn Thr Thr His Ile Trp Tyr Thr
 65 70 75 80
 Cys His Met Arg Leu Ser Gln Phe Leu Ser Asp Glu Val Phe Ile Val
 85 90 95
 Asn Val Thr Asp Gln Ser Gly Asn Asn Ser Gln Glu Cys Gly Ser Phe
 100 105 110
 Val Leu Ala Glu Ser Ile Lys Pro Ala Pro Pro Leu Asn Val Thr Val
 115 120 125
 Ala Phe Ser Gly Arg Tyr Asp Ile Ser Trp Asp Ser Ala Tyr Asp Glu
 130 135 140
 Pro Ser Asn Tyr Val Leu Arg Gly Lys Leu Gln Tyr Glu Leu Gln Tyr
 145 150 155 160
 Arg Asn Leu Arg Asp Pro Tyr Ala Val Arg Pro Val Thr Lys Leu Ile
 165 170 175
 Ser Val Asp Ser Arg Asn Val Ser Leu Leu Pro Glu Glu Phe His Lys
 180 185 190
 Asp Ser Ser Tyr Gln Leu Gln Val Arg Ala Ala Pro Gln Pro Gly Thr
 195 200 205
 Ser Phe Arg Gly Thr Trp Ser Glu Trp Ser Asp Pro Val Ile Phe Gln
 210 215 220
 Thr Gln Ala Gly Glu Pro Glu Ala Gly Trp Asp Pro His Met Leu Leu
 225 230 235 240
 Leu Leu Ala Val Leu Ile Ile Val Leu Val Phe Met Gly Leu Lys Ile
 245 250 255
 His Leu Pro Trp Arg Leu Trp Lys Lys Ile Trp Ala Pro Val Pro Thr
 260 265 270
 Pro Glu Ser Phe Phe Gln Pro Leu Tyr Arg Glu His Ser Gly Asn Phe
 275 280 285
 Lys Lys Trp Val Asn Thr Pro Phe Thr Ala Ser Ser Ile Glu Leu Val
 290 295 300
 Pro Gln Ser Ser Thr Thr Thr Ser Ala Leu His Leu Ser Leu Tyr Pro
 305 310 315 320
 Ala Lys Glu Lys Lys Phe Pro Gly Leu Pro Gly Leu Glu Glu Gln Leu
 325 330 335

Glu Cys Asp Gly Met Ser Glu Pro Gly His Trp Cys Ile Ile Pro Leu
 340 345 350
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 355 360 365
 Tyr Gly Leu Val Ser Ile Asp Thr Val Thr Val Gly Asp Ala Glu Gly
 370 375 380
 Leu Cys Val Trp Pro Cys Ser Cys Glu Asp Asp Gly Tyr Pro Ala Met
 385 390 395 400
 Asn Leu Asp Ala Gly Arg Glu Ser Gly Pro Asn Ser Glu Asp Leu Leu
 405 410 415
 Leu Val Thr Asp Pro Ala Phe Leu Ser Cys Gly Cys Val Ser Gly Ser
 420 425 430
 Gly Leu Arg Leu Gly Gly Ser Pro Gly Ser Leu Leu Asp Arg Leu Arg
 435 440 445
 Leu Ser Phe Ala Lys Glu Gly Asp Trp Thr Ala Asp Pro Thr Trp Arg
 450 455 460
 Thr Gly Ser Pro Gly Gly Gly Ser Glu Ser Glu Ala Gly Ser Pro Pro
 465 470 475 480
 Gly Leu Asp Met Asp Thr Phe Asp Ser Gly Phe Ala Gly Ser Asp Cys
 485 490 495
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 500 505 510
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 515 520 525
 Ser

<210> 11
 <211> 20
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Description of Artificial Sequence: PCR Primer

<400> 11
 agcatcaagc cggctccccc

20

<210> 12
 <211> 20
 <212> DNA
 <213> Artificial Primer

<220>
 <223> Description of Artificial Sequence: PCR Primer

<400> 12
 ctccattcac tccaggtccc

20

<210> 13
 <211> 20
 <212> DNA
 <213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: PCR Primer

<400> 13

ttgaacgtga ctgrggcctt

20

<210> 14

<211> 20

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: Murine MU-1 cDNA Internal Oligonucleotide

<400> 14

tgaatgaagt gcctggctga

20

<210> 15

<211> 40

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: 5' PCR Primer

<400> 15

cacaaagctt cagtatgagc tgcagtacag gaaccgggga

40

<210> 16

<211> 40

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: 3' PCR primer

<400> 16

cacaggatcc ctttaactcc tctgactggg tctgaaagat

40

<210> 17

<211> 224

<212> PRT

<213> Unknown Organism

<220>

<221> Unknown Organism

<222> (1)..(224)

<223> Description of Unknown Organism: Second polypeptide comprising an Fc region

<400> 17

His	Thr	Cys	Pro	Pro	Cys	Pro	Ala	Pro	Glu	Ala	Leu	Gly	Ala	Pro	Ser
1				5					10					15	

Val	Phe	Leu	Phe	Pro	Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg
		20						25					30		

Thr	Pro	Glu	Val	Thr	Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro
		35					40					45			

Glu	Val	Lys	Phe	Asn	Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala
	50					55					60				

Lys Thr Lys Pro Arg Glu Glu Gln Tyr Asn Ser Thr Tyr Arg Val Val
 65 70 75 80
 Ser Val Leu Thr Val Leu His Gln Asp Trp Leu Asn Gly Lys Glu Tyr
 85 90 95
 Lys Cys Lys Val Ser Asn Lys Ala Leu Pro Val Pro Ile Glu Lys Thr
 100 105 110
 Ile Ser Lys Ala Lys Gly Gln Pro Arg Glu Pro Gln Val Tyr Thr Leu
 115 120 125
 Pro Pro Ser Arg Glu Glu Met Thr Lys Asn Gln Val Ser Leu Thr Cys
 130 135 140
 Leu Val Lys Gly Phe Tyr Pro Ser Asp Ile Ala Val Glu Trp Glu Ser
 145 150 155 160
 Asn Gly Gln Pro Glu Asn Asn Tyr Lys Thr Thr Pro Pro Val Leu Asp
 165 170 175
 Ser Asp Gly Ser Phe Phe Leu Tyr Ser Lys Leu Thr Val Asp Lys Ser
 180 185 190
 Arg Trp Gln Gln Gly Asn Val Phe Ser Cys Ser Val Met His Glu Ala
 195 200 205
 Leu His Asn His Tyr Thr Gln Lys Ser Leu Ser Leu Ser Pro Gly Lys
 210 215 220

<210> 18
 <211> 617
 <212> DNA
 <213> Human

<400> 18
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 tggcaacatg gagaggattg tcatctgtct gatggtcac ttcttgggga cactgggtcca 120
 caaatcaagc tccaagggtc aagatcgcca catgattaga atgcgtcaac ttatagatat 180
 tgttgatcag ctgaaaaatt atgtgaatga cttgggtccct gaatttctgc cagctccaga 240
 agatgtagag acaaactgtg agtgggtcagc ttttcctgc tttcagaagg cccaactaaa 300
 gtcagcaaat acaggaaaca atgaaaggat aatcaatgta tcaattaaaa agctgaagag 360
 gaaaccacct tccacaaatg cagggagaag acagaaacac agactaacat gcccttcacg 420
 tgattcttat gagaaaaaac catccaaaga attcctagaa agattcaaatt cacttctcca 480
 aaagatgatt catcagcatc tgtcctctag aacacacgga agtgaagatt cctgaggatc 540
 taacttgcag ttggacacta tggtacatac tctaatatag tagtgaaagt catttctttg 600
 tattccaagt ggaggag 617

<210> 19
 <211> 162
 <212> PRT
 <213> Human

<400> 19

Met Arg Ser Ser Pro Gly Asn Met Glu Arg Ile Val Ile Cys Leu Met
 1 5 10 15

Val Ile Phe Leu Gly Thr Leu Val His Lys Ser Ser Ser Gln Gly Gln
 20 25 30
 Asp Arg His Met Ile Arg Met Arg Gln Leu Ile Asp Ile Val Asp Gln
 35 40 45
 Leu Lys Asn Tyr Val Asn Asp Leu Val Pro Glu Phe Leu Pro Ala Pro
 50 55 60
 Glu Asp Val Glu Thr Asn Cys Glu Trp Ser Ala Phe Ser Cys Phe Gln
 65 70 75 80
 Lys Ala Gln Leu Lys Ser Ala Asn Thr Gly Asn Asn Glu Arg Ile Ile
 85 90 95
 Asn Val Ser Ile Lys Lys Leu Lys Arg Lys Pro Pro Ser Thr Asn Ala
 100 105 110
 Gly Arg Arg Gln Lys His Arg Leu Thr Cys Pro Ser Cys Asp Ser Tyr
 115 120 125
 Glu Lys Lys Pro Pro Lys Glu Phe Leu Glu Arg Phe Lys Ser Leu Leu
 130 135 140
 Gln Lys Met Ile His Gln His Leu Ser Ser Arg Thr His Gly Ser Glu
 145 150 155 160
 Asp Ser

<210> 20
 <211> 7
 <212> PRT
 <213> Human

<400> 20

Glu Asp Asp Gly Tyr Pro Ala
1 5

<210> 21
 <211> 16
 <212> PRT
 <213> Human

<400> 21

Met Pro Leu Leu Leu Leu Leu Leu Leu Pro Ser Pro Leu His Pro
1 5 10 15

<210> 22
 <211> 786
 <212> DNA
 <213> Human

<400> 22
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 gccggcagcg gacaccacca tcatcaccac ggtagcggcg actataaaga cgatgacgat 120
 aagggttccg gatgccccga cctcgctgc tacaccgatt acctccagac ggtcatctgc 180
 atcctggaaa tgtggaacct ccaccccagc acgctcacc ttacctggca agaccagtat 240
 gaagagctga aggacgaggc cacctcctgc agcctccaca ggtcggccca caatgccacg 300

catgccacct acacctgcca catggatgta ttccacttca tggccgacga cattttcagt 360
gtcaacatca cagaccagtc tggcaactac tcccaggagt gtggcagctt tctcctggct 420
gagagcatca agccggctcc ccctttcaac gtgactgtga ccttctcagg acagtataat 480
atctcctggc gctcagatta cgaagaccct gccttctaca tgctgaaggg caagcttcag 540
tatgagctgc agtacaggaa ccggggagac ccctgggctg tgagtccgag gagaaagctg 600
atctcagtggt actcaagaag tgtctccctc ctccccctgg agttccgcaa agactcgagc 660
tatgagctgc aggtgcgggc agggcccatg cctggctcct cctaccaggg gacctggagt 720
gaatggagtg acccggtcat ctttcagacc cagtcagagg agttaaagga aggctggaac 780
taatga 786

<210> 23
<211> 260
<212> PRT
<213> Human

<400> 23

Met Lys Phe Leu Val Asn Val Ala Leu Val Phe Met Val Val Tyr Ile
1 5 10 15
Ser Tyr Ile Tyr Ala Gly Ser Gly His His His His His His Gly Ser
20 25 30
Gly Asp Tyr Lys Asp Asp Asp Asp Lys Gly Ser Gly Cys Pro Asp Leu
35 40 45
Val Cys Tyr Thr Asp Tyr Leu Gln Thr Val Ile Cys Ile Leu Glu Met
50 55 60
Trp Asn Leu His Pro Ser Thr Leu Thr Leu Thr Trp Gln Asp Gln Tyr
65 70 75 80
Glu Glu Leu Lys Asp Glu Ala Thr Ser Cys Ser Leu His Arg Ser Ala
85 90 95
His Asn Ala Thr His Ala Thr Tyr Thr Cys His Met Asp Val Phe His
100 105 110
Phe Met Ala Asp Asp Ile Phe Ser Val Asn Ile Thr Asp Gln Ser Gly
115 120 125
Asn Tyr Ser Gln Glu Cys Gly Ser Phe Leu Leu Ala Glu Ser Ile Lys
130 135 140
Pro Ala Pro Pro Phe Asn Val Thr Val Thr Phe Ser Gly Gln Tyr Asn
145 150 155 160
Ile Ser Trp Arg Ser Asp Tyr Glu Asp Pro Ala Phe Tyr Met Leu Lys
165 170 175
Gly Lys Leu Gln Tyr Glu Leu Gln Tyr Arg Asn Arg Gly Asp Pro Trp
180 185 190
Ala Val Ser Pro Arg Arg Lys Leu Ile Ser Val Asp Ser Arg Ser Val
195 200 205
Ser Leu Leu Pro Leu Glu Phe Arg Lys Asp Ser Ser Tyr Glu Leu Gln
210 215 220

Val Arg Ala Gly Pro Met Pro Gly Ser Ser Tyr Gln Gly Thr Trp Ser
 225 230 235 240
 Glu Trp Ser Asp Pro Val Ile Phe Gln Thr Gln Ser Glu Glu Leu Lys
 245 250 255
 Glu Gly Trp Asn
 260

<210> 24
 <211> 1426
 <212> DNA
 <213> Human

<400> 24
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 gaggctgggg ctgccccgac ctgctctgct acaccgatta cctccagacg gtcctctgca 120
 tcctggaaat gtggaacctc caccgccagca cgctcaccct tacctggcaa gaccagtatg 180
 aagagctgaa ggacgaggcc acctcctgca gcctccacag gtcggccac aatgccacgc 240
 atgccacctc cacctgccac atggatgtat tccattcat ggccgacgac attttcagtg 300
 tcaacatcac agaccagtct ggcaactact ccaggagtg tggcagcttt ctctggctg 360
 agagcatcaa gccggctccc cttttcaacg tgactgtgac cttctcagga cagtataata 420
 tctcctggcg ctgagattac gaagaccctg cttctacat gctgaagggc aagcttcagt 480
 atgagctgca gtacaggaac cggggagacc cctgggctgt gagtccgagg agaaagctga 540
 tctcagtggc ctcaagaagt gtctccctcc tccccctgga gttccgcaaa gactcgagct 600
 atgagctgca ggtgcgggca gggcccatgc ctggctcctc ctaccagggg acctggagtg 660
 aatggagtga cccggtcatc tttcagaccc agtcagagga gttaaaggaa ggctggaacg 720
 gtcctggctc tagagacaaa actcacacat gccaccctg cccagcacct gaactcctgg 780
 ggggaccgtc agtcttcctc tccccccaa aaccaagga caccctcatg atctcccgga 840
 cccctgaggt cacatgctg gtggtggacg tgagccacga agaccctgag gtcaagttca 900
 actggtacgt ggacggcgtg gaggtgcata atgccaagac aaagccgcgg gaggagcagt 960
 acaacagcac gtaccgtgtg gtcagcgtcc tcaccgtcct gcaccaggac tggctgaatg 1020
 gcaaggagta caagtgaag gtctccaaca aagccctccc agtccccatc gagaaaacca 1080
 tctccaaagc caaagggcag ccccgagaac cacaggtgta caccctgccc ccatcccgga 1140
 aggagatgac caagaaccag gtcagcctga cctgcctggg caaaggcttc tatcccagcg 1200
 acatcgccgt ggagtgggag agcaatgggc agccggagaa caactacaag accacgcctc 1260
 ccgtgctgga ctccgacggc tccttcttcc tctatagcaa gtcaccgtg gacaagagca 1320
 ggtggcagca ggggaacgtc ttctcatgct ccgtgatgca tgaggctctg cacaaccact 1380
 acacgcagaa gagcctctcc ctgtccccgg gtaaatgagt gaattc 1426

<210> 25
 <211> 467
 <212> PRT
 <213> Human

<400> 25

Met Pro Arg Gly Trp Ala Ala Pro Leu Leu Leu Leu Leu Leu Gln Gly
 1 5 10 15
 Gly Trp Gly Cys Pro Asp Leu Val Cys Tyr Thr Asp Tyr Leu Gln Thr
 20 25 30
 Val Ile Cys Ile Leu Glu Met Trp Asn Leu His Pro Ser Thr Leu Thr
 35 40 45
 Leu Thr Trp Gln Asp Gln Tyr Glu Glu Leu Lys Asp Glu Ala Thr Ser
 50 55 60
 Cys Ser Leu His Arg Ser Ala His Asn Ala Thr His Ala Thr Tyr Thr
 65 70 75 80
 Cys His Met Asp Val Phe His Phe Met Ala Asp Asp Ile Phe Ser Val
 85 90 95
 Asn Ile Thr Asp Gln Ser Gly Asn Tyr Ser Gln Glu Cys Gly Ser Phe
 100 105 110
 Leu Leu Ala Glu Ser Ile Lys Pro Ala Pro Pro Phe Asn Val Thr Val
 115 120 125
 Thr Phe Ser Gly Gln Tyr Asn Ile Ser Trp Arg Ser Asp Tyr Glu Asp
 130 135 140
 Pro Ala Phe Tyr Met Leu Lys Gly Lys Leu Gln Tyr Glu Leu Gln Tyr
 145 150 155 160
 Arg Asn Arg Gly Asp Pro Trp Ala Val Ser Pro Arg Arg Lys Leu Ile
 165 170 175
 Ser Val Asp Ser Arg Ser Val Ser Leu Leu Pro Leu Glu Phe Arg Lys
 180 185 190
 Asp Ser Ser Tyr Glu Leu Gln Val Arg Ala Gly Pro Met Pro Gly Ser
 195 200 205
 Ser Tyr Gln Gly Thr Trp Ser Glu Trp Ser Asp Pro Val Ile Phe Gln
 210 215 220
 Thr Gln Ser Glu Glu Leu Lys Glu Gly Trp Asn Gly Ser Gly Ser Arg
 225 230 235 240
 Asp Lys Thr His Thr Cys Pro Pro Cys Pro Ala Pro Glu Leu Leu Gly
 245 250 255
 Gly Pro Ser Val Phe Leu Phe Pro Pro Lys Pro Lys Asp Thr Leu Met
 260 265 270
 Ile Ser Arg Thr Pro Glu Val Thr Cys Val Val Val Asp Val Ser His
 275 280 285
 Glu Asp Pro Glu Val Lys Phe Asn Trp Tyr Val Asp Gly Val Glu Val
 290 295 300
 His Asn Ala Lys Thr Lys Pro Arg Glu Glu Gln Tyr Asn Ser Thr Tyr
 305 310 315 320
 Arg Val Val Ser Val Leu Thr Val Leu His Gln Asp Trp Leu Asn Gly
 325 330 335
 Lys Glu Tyr Lys Cys Lys Val Ser Asn Lys Ala Leu Pro Val Pro Ile
 340 345 350

Glu Lys Thr Ile Ser Lys Ala Lys Gly Gln Pro Arg Glu Pro Gln Val
 355 360 365
 Tyr Thr Leu Pro Pro Ser Arg Glu Glu Met Thr Lys Asn Gln Val Ser
 370 375 380
 Leu Thr Cys Leu Val Lys Gly Phe Tyr Pro Ser Asp Ile Ala Val Glu
 385 390 395 400
 Trp Glu Ser Asn Gly Gln Pro Glu Asn Asn Tyr Lys Thr Thr Pro Pro
 405 410 415
 Val Leu Asp Ser Asp Gly Ser Phe Phe Leu Tyr Ser Lys Leu Thr Val
 420 425 430
 Asp Lys Ser Arg Trp Gln Gln Gly Asn Val Phe Ser Cys Ser Val Met
 435 440 445
 His Glu Ala Leu His Asn His Tyr Thr Gln Lys Ser Leu Ser Leu Ser
 450 455 460
 Pro Gly Lys
 465

<210> 26
 <211> 1499
 <212> DNA
 <213> Human

<400> 26
 gcggccgcac caccatgccg cgtggctggg ccgccccctt gtcctgctg ctgctccagg 60
 gagggctggg ctgccccgac ctgctctgct acaccgatta cctccagacg gtcattctgca 120
 tcctggaaat gtggaacctc caccacagca cgctcaccct tacctggcaa gaccagtatg 180
 aagagctgaa ggacgaggcc acctcctgca gcctccacag gtcggccac aatgccacgc 240
 atgccaccta cacctgccac atggatgtat tccacttcat ggccgacgac attttcagtg 300
 tcaacatcac agaccagtct ggcaactact cccaggagtg tggcagcttt ctctgggctg 360
 agagcatcaa gccggctccc cttttcaacg tgactgtgac cttctcagga cagtataata 420
 tctcctggcg ctgagattac gaagaccctg ctttctacat gctgaagggc aagcttcagt 480
 atgagctgca gtacaggaac cggggagacc cctgggctgt gaggccgagg agaaagctga 540
 tctcagtggg ctcaagaagt gtctccctcc tccccctgga gtcccgcaaa gactcgagct 600
 atgagctgca ggtgcgggca gggcccatgc ctggctcctc ctaccagggg acctggagtg 660
 aatggagtga cccggctatc ttccagaccc agtcagagga gttaaaggaa ggctggaacg 720
 gtcctggctc tagagacaaa actcacacat gccacccgtg cccagcacct gaactcctgg 780
 ggggaccgtc agtcttcctc tccccccaa aacccaagga caccctcatg atctcccgga 840
 cccctgaggt cacatgcgtg gtgggtggacg tgagccacga agaccctgag gtcaagttca 900
 actggtacgt ggacggcgtg gaggtgcata atgccaagac aaagccgcgg gaggagcagt 960
 acaacagcac gtaccgtgtg gtcagcgtcc tcaccgtcct gcaccaggac tggctgaatg 1020
 gcaaggagta caagtgaag gtctccaaca aagccctccc agtccccatc gagaaaacca 1080
 tctcaaagc caaagggcag ccccgagaac cacaggtgta caccctgccc ccatcccggg 1140

aggagatgac caagaaccag gtcagcctga cctgcctggt caaaggcttc tatcccagcg 1200
 acatcgccgt ggagtgggag agcaatgggc agccggagaa caactacaag accacgcctc 1260
 ccgtgctgga ctccgacggc tccttcttcc tctatagcaa gctcaccgtg gacaagagca 1320
 ggtggcagca ggggaacgtc ttctcatgct ccgtgatgca tgaggctctg cacaaccact 1380
 acacgcagaa gagcctctcc ctgtccccgg gtaaattcagg aatggcatca atgacaggag 1440
 gtcaacaaat gggttctgga tctcatcatc atcatcatca ttctggaggt tgagaattc 1499

<210> 27
 <211> 492
 <212> PRT
 <213> Human

<400> 27

Met Pro Arg Gly Trp Ala Ala Pro Leu Leu Leu Leu Leu Gln Gly
 1 5 10 15
 Gly Trp Gly Cys Pro Asp Leu Val Cys Tyr Thr Asp Tyr Leu Gln Thr
 20 25 30
 Val Ile Cys Ile Leu Glu Met Trp Asn Leu His Pro Ser Thr Leu Thr
 35 40 45
 Leu Thr Trp Gln Asp Gln Tyr Glu Glu Leu Lys Asp Glu Ala Thr Ser
 50 55 60
 Cys Ser Leu His Arg Ser Ala His Asn Ala Thr His Ala Thr Tyr Thr
 65 70 75 80
 Cys His Met Asp Val Phe His Phe Met Ala Asp Asp Ile Phe Ser Val
 85 90 95
 Asn Ile Thr Asp Gln Ser Gly Asn Tyr Ser Gln Glu Cys Gly Ser Phe
 100 105 110
 Leu Leu Ala Glu Ser Ile Lys Pro Ala Pro Pro Phe Asn Val Thr Val
 115 120 125
 Thr Phe Ser Gly Gln Tyr Asn Ile Ser Trp Arg Ser Asp Tyr Glu Asp
 130 135 140
 Pro Ala Phe Tyr Met Leu Lys Gly Lys Leu Gln Tyr Glu Leu Gln Tyr
 145 150 155 160
 Arg Asn Arg Gly Asp Pro Trp Ala Val Ser Pro Arg Arg Lys Leu Ile
 165 170 175
 Ser Val Asp Ser Arg Ser Val Ser Leu Leu Pro Leu Glu Phe Arg Lys
 180 185 190
 Asp Ser Ser Tyr Glu Leu Gln Val Arg Ala Gly Pro Met Pro Gly Ser
 195 200 205
 Ser Tyr Gln Gly Thr Trp Ser Glu Trp Ser Asp Pro Val Ile Phe Gln
 210 215 220
 Thr Gln Ser Glu Glu Leu Lys Glu Gly Trp Asn Gly Ser Gly Ser Arg
 225 230 235 240
 Asp Lys Thr His Thr Cys Pro Pro Cys Pro Ala Pro Glu Leu Leu Gly
 245 250 255

Gly Pro Ser Val Phe Leu Phe Pro Pro Lys Pro Lys Asp Thr Leu Met
 260 265 270
 Ile Ser Arg Thr Pro Glu Val Thr Cys Val Val Val Asp Val Ser His
 275 280 285
 Glu Asp Pro Glu Val Lys Phe Asn Trp Tyr Val Asp Gly Val Glu Val
 290 295 300
 His Asn Ala Lys Thr Lys Pro Arg Glu Glu Gln Tyr Asn Ser Thr Tyr
 305 310 315 320
 Arg Val Val Ser Val Leu Thr Val Leu His Gln Asp Trp Leu Asn Gly
 325 330 335
 Lys Glu Tyr Lys Cys Lys Val Ser Asn Lys Ala Leu Pro Val Pro Ile
 340 345 350
 Glu Lys Thr Ile Ser Lys Ala Lys Gly Gln Pro Arg Glu Pro Gln Val
 355 360 365
 Tyr Thr Leu Pro Pro Ser Arg Glu Glu Met Thr Lys Asn Gln Val Ser
 370 375 380
 Leu Thr Cys Leu Val Lys Gly Phe Tyr Pro Ser Asp Ile Ala Val Glu
 385 390 395 400
 Trp Glu Ser Asn Gly Gln Pro Glu Asn Asn Tyr Lys Thr Thr Pro Pro
 405 410 415
 Val Leu Asp Ser Asp Gly Ser Phe Phe Leu Tyr Ser Lys Leu Thr Val
 420 425 430
 Asp Lys Ser Arg Trp Gln Gln Gly Asn Val Phe Ser Cys Ser Val Met
 435 440 445
 His Glu Ala Leu His Asn His Tyr Thr Gln Lys Ser Leu Ser Leu Ser
 450 455 460
 Pro Gly Lys Ser Gly Met Ala Ser Met Thr Gly Gly Gln Gln Met Gly
 465 470 475 480
 Ser Gly Ser His His His His His His Ser Gly Gly
 485 490

<210> 28
 <211> 1426
 <212> DNA
 <213> Human

<400> 28
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 gaggctgggg ctgccccgac ctgctctgct acaccgatta cctccagacg gtcattctgca 120
 tcctggaaat gtggaacctc caccacagca cgctcaccct tacctggcaa gaccagtatg 180
 aagagctgaa ggacgaggcc acctcctgca gcctccacag gtcggccac aatgccacgc 240
 atgccaccta cacctgccac atggatgtat tccacttcat ggccgacgac attttcagt 300
 tcaacatcac agaccagtct ggcaactact cccaggagtg tggcagcttt ctcctggctg 360
 agagcatcaa gccggctccc cttttcaacg tgactgtgac cttctcagga cagtataata 420
 tctcctggcg ctgagattac gaagaacctg cttctacat gctgaagggc aagcttcagt 480
 atgagctgca gtacaggaac cggggagacc cctgggctgt gaggccgagg agaaagctga 540

tctcagtgga ctcaagaagt gtctccctcc tccccctgga gttccgcaaa gactcgagct 600
 atgagctgca ggtgcgggca gggcccatgc ctggctcctc ctaccagggg acctggagtg 660
 aatggagtga cccggtcatc ttccagaccc agtcagagga gttaaaggaa ggctggaacg 720
 gctccggctc tagagacaaa actcacacat gcccaccgtg cccagcacct gaagccctgg 780
 gggcaccgtc agtcttcctc ttcccccaa aaccaagga caccctcatg atctcccgga 840
 cccctgaggt cacatgcgtg gtggtggacg tgagccacga agaccctgag gtcaagttca 900
 actggtacgt ggacggcgtg gaggtgcata atgccaagac aaagccgcgg gaggagcagt 960
 acaacagcac gtaccgtgtg gtcagcgtcc tcaccgtcct gcaccaggac tggctgaatg 1020
 gcaaggagta caagtgaag gtctccaaca aagccctccc agcccccatc gagaaaacca 1080
 tctccaaagc caaagggcag ccccgagaac cacagggtga caccctgccc ccatcccggg 1140
 aggagatgac caagaaccag gtcagcctga cctgcctggg caaaggcttc tatcccagcg 1200
 acatcgccgt ggagtgggag agcaatgggc agccggagaa caactacaag accacgcctc 1260
 ccgtgctgga ctccgacggc tccttcttcc tctatagcaa gtcaccgtg gacaagagca 1320
 ggtggcagca ggggaacgtc ttctcatgct ccgtgatgca tgaggctctg cacaaccact 1380
 acacgcagaa gagcctctcc ctgtccccgg gtaaattgagt gaattc 1426

<210> 29
 <211> 467
 <212> PRT
 <213> Human

<400> 29

Met Pro Arg Gly Trp Ala Ala Pro Leu Leu Leu Leu Leu Leu Gln Gly
 1 5 10 15
 Gly Trp Gly Cys Pro Asp Leu Val Cys Tyr Thr Asp Tyr Leu Gln Thr
 20 25 30
 Val Ile Cys Ile Leu Glu Met Trp Asn Leu His Pro Ser Thr Leu Thr
 35 40 45
 Leu Thr Trp Gln Asp Gln Tyr Glu Glu Leu Lys Asp Glu Ala Thr Ser
 50 55 60
 Cys Ser Leu His Arg Ser Ala His Asn Ala Thr His Ala Thr Tyr Thr
 65 70 75 80
 Cys His Met Asp Val Phe His Phe Met Ala Asp Asp Ile Phe Ser Val
 85 90 95
 Asn Ile Thr Asp Gln Ser Gly Asn Tyr Ser Gln Glu Cys Gly Ser Phe
 100 105 110
 Leu Leu Ala Glu Ser Ile Lys Pro Ala Pro Pro Phe Asn Val Thr Val
 115 120 125
 Thr Phe Ser Gly Gln Tyr Asn Ile Ser Trp Arg Ser Asp Tyr Glu Asp
 130 135 140
 Pro Ala Phe Tyr Met Leu Lys Gly Lys Leu Gln Tyr Glu Leu Gln Tyr
 145 150 155 160

Arg Asn Arg Gly Asp Pro Trp Ala Val Ser Pro Arg Arg Lys Leu Ile
 165 170 175
 Ser Val Asp Ser Arg Ser Val Ser Leu Leu Pro Leu Glu Phe Arg Lys
 180 185 190
 Asp Ser Ser Tyr Glu Leu Gln Val Arg Ala Gly Pro Met Pro Gly Ser
 195 200 205
 Ser Tyr Gln Gly Thr Trp Ser Glu Trp Ser Asp Pro Val Ile Phe Gln
 210 215 220
 Thr Gln Ser Glu Glu Leu Lys Glu Gly Trp Asn Gly Ser Gly Ser Arg
 225 230 235 240
 Asp Lys Thr His Thr Cys Pro Pro Cys Pro Ala Pro Glu Ala Leu Gly
 245 250 255
 Ala Pro Ser Val Phe Leu Phe Pro Pro Lys Pro Lys Asp Thr Leu Met
 260 265 270
 Ile Ser Arg Thr Pro Glu Val Thr Cys Val Val Val Asp Val Ser His
 275 280 285
 Glu Asp Pro Glu Val Lys Phe Asn Trp Tyr Val Asp Gly Val Glu Val
 290 295 300
 His Asn Ala Lys Thr Lys Pro Arg Glu Glu Gln Tyr Asn Ser Thr Tyr
 305 310 315 320
 Arg Val Val Ser Val Leu Thr Val Leu His Gln Asp Trp Leu Asn Gly
 325 330 335
 Lys Glu Tyr Lys Cys Lys Val Ser Asn Lys Ala Leu Pro Ala Pro Ile
 340 345 350
 Glu Lys Thr Ile Ser Lys Ala Lys Gly Gln Pro Arg Glu Pro Gln Val
 355 360 365
 Tyr Thr Leu Pro Pro Ser Arg Glu Glu Met Thr Lys Asn Gln Val Ser
 370 375 380
 Leu Thr Cys Leu Val Lys Gly Phe Tyr Pro Ser Asp Ile Ala Val Glu
 385 390 395 400
 Trp Glu Ser Asn Gly Gln Pro Glu Asn Asn Tyr Lys Thr Thr Pro Pro
 405 410 415
 Val Leu Asp Ser Asp Gly Ser Phe Phe Leu Tyr Ser Lys Leu Thr Val
 420 425 430
 Asp Lys Ser Arg Trp Gln Gln Gly Asn Val Phe Ser Cys Ser Val Met
 435 440 445
 His Glu Ala Leu His Asn His Tyr Thr Gln Lys Ser Leu Ser Leu Ser
 450 455 460
 Pro Gly Lys
 465

<210> 30
 <211> 741
 <212> DNA
 <213> Human

<400> 30

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cccgacctcg tctgctacac cgattacctc cagacgggtca tctgcatcct ggaaatgtgg      120
aacctccacc ccagcacgct cacccttacc tggcaagacc agtatgaaga gctgaaggac      180
gaggccacct cctgcagcct ccacaggtcg gcccacaatg ccacgcatgc cacctacacc      240
tgccacatgg atgtattcca cttcatggcc gacgacattt tcagtgtcaa catcacagac      300
cagtctggca actactccca ggagtgtggc agcttttctcc tggctgagag catcaagccg      360
gctccccctt tcaacgtgac tgtgaccttc tcaggacagt ataatatctc ctggcgctca      420
gattacgaag accctgcctt ctacatgctg aagggcaagc ttcagtatga gctgcagtag      480
aggaaccggg gagaccctcg ggctgtgagt ccgaggagaa agctgatctc agtggactca      540
agaagtgtct ccctcctccc cctggagttc cgcaaagact cgagctatga gctgcaggtg      600
cgggcagggc ccatgcctgg ctctcctac caggggacct ggagtgaatg gagtgacctg      660
gtcatctttc agaccagtc agaggagtta aaggaaggct ggaacaaaac cgaaacctcc      720
caggttgctc cggcataatg a                                         741
  
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<210> 31
 <211> 245
 <212> PRT
 <213> Human

<400> 31

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Met Pro Arg Gly Trp Ala Ala Pro Leu Leu Leu Leu Leu Gln Gly
1          5          10          15
Gly Trp Gly Cys Pro Asp Leu Val Cys Tyr Thr Asp Tyr Leu Gln Thr
20          25          30
Val Ile Cys Ile Leu Glu Met Trp Asn Leu His Pro Ser Thr Leu Thr
35          40          45
Leu Thr Trp Gln Asp Gln Tyr Glu Glu Leu Lys Asp Glu Ala Thr Ser
50          55          60
Cys Ser Leu His Arg Ser Ala His Asn Ala Thr His Ala Thr Tyr Thr
65          70          75          80
Cys His Met Asp Val Phe His Phe Met Ala Asp Asp Ile Phe Ser Val
85          90          95
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 195 200 205
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 50 55 60
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 Cys His Met Asp Val Phe His Phe Met Ala Asp Asp Ile Phe Ser Val
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 Pro Ser Asn Tyr Val Leu Arg Gly Lys Leu Gln Tyr Glu Leu Gln Tyr
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Arg Asn Leu Arg Asp Pro Tyr Ala Val Arg Pro Val Thr Lys Leu Ile
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 His Asn Thr Thr His Ile Trp Tyr Thr Cys His Met Arg Leu Ser Gln
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 195 200 205
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 210 215 220
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<400> 41

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<223> Description of Artificial Sequence: 5' PCR Primer

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<212> DNA
<213> Artificial Sequence

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<223> Description of Artificial Sequence: 5' PCR Primer

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<212> DNA
<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: PCR Probe

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26

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GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW,
MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG,
SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ,
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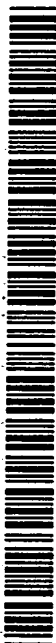
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(54) Title: METHODS AND COMPOSITIONS FOR MODULATING INTERLEUKIN-21 RECEPTOR ACTIVITY

(57) Abstract: Methods and compositions for modulating interleukin-21 (IL-21)/IL-21 receptor (MU-1) activity using agonists or antagonists of IL-21 or IL-21 receptor ("IL-21R" or "MU-1"), are disclosed. IL-21/IL-21R antagonists can be used to induce immune suppression *in vivo*, e.g., for treating or preventing immune cell-associated pathologies (e.g., pathologies associated with aberrant activity of one or more of mature T cells (mature CD8+, mature CD4+ T cells), mature NK cells, B cells, macrophages and megakaryocytes, including transplant rejection and autoimmune disorders). IL-21/IL-21R agonists can be used by themselves or in combination with an antigen, e.g., as an adjuvant (e.g., a vaccine adjuvant), to up-regulate an immune response *in vivo*, e.g., for example, for use in treating cancer and infectious disorders.



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International application No.

PCT/US02/29839

A. CLASSIFICATION OF SUBJECT MATTER

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According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 514/2, 12; 530/350, 387.3; 435/69.1, 252.3, 357, 254.11, 320.1; 424/134.1, 192.1

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
Commercial and Patent Sequence Databases: SEQ ID NO: 2

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X — Y	US 6,057,128 A (DONALDSON et al) 2 May 2000 (02.05.00), SEQ ID NO: 2, column, 3, lines 49-54, column 4, line 12 to column 5, line 8, column 6, line 1-31, column 6, line 46 to column 8, line 53, column 9, line 44-58, column 10, line 56 to column 11, line 11 and column 16, lines 19-24.	1, 10, 11, 13, 14, 37-41, 45-47, 2-9, 12, 42-44, 48-60
X — Y	WO 00/17235 A2 (ZYMOGENETICS, INC.) 30 May 2000(30.03.00), SEQ ID NO: 2, pages 2-5, 7, 39-46, 52-54, 66 and 69.	1, 10, 11, 13, 14, 37-41, 45-47 2-9, 12, 42-44, 48-60
X — Y	WO 00/27882 A1 (SMITHKLINE BEECHAM CORPORATION) 18 May 2000 (18.03.00), SEQ ID NO: 2, pages 2, 6-8, 15 and claim 4.	1, 10, 11, 13, 14 2-9, 12, 37-60
T	US 2002/0137677 A1 (SPRECHER et al.) 26 September 2002 (26.09.02).	1-14, 37-60

☐ Further documents are listed in the continuation of Box C.

☐ See patent family annex.

Special categories of cited documents:	
A document defining the general state of the art which is not considered to be of particular relevance	*T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
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O document referring to an oral disclosure, use, exhibition or other means	*&* document member of the same patent family
P document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

13 November 2003 (13.11.2003)

Date of mailing of the international search report

18 DEC 2003

Name and mailing address of the ISA/US

Mail Stop PCT, Attn: ISA/US
Commissioner for Patents
P.O. Box 1450
Alexandria, Virginia 22313-1450

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INTERNATIONAL SEARCH REPORT

International application No.

PCT/US02/29839

Box I Observations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)

This international report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claim Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☐ Claim Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☒ Claim Nos.: 15-23,36 and 61-63
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of Item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:
Please See Continuation Sheet

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.: 1-14 and 37-60

Remark on Protest

☐
☐

The additional search fees were accompanied by the applicant's protest.

No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

PCT/US02/29839

BOX II. OBSERVATIONS WHERE UNITY OF INVENTION IS LACKING

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1. In order for all inventions to be examined, the appropriate additional examination fees must be paid.

Group I, claim(s) 1-14 and 37-60, in so far as they are drawn to IL-21/IL-21R antagonist that is fragment of human IL-21 receptor of fusion protein thereof with Ig, method of treating or preventing an arthritic disorder and method of modulating the activity of an immune cell. Within Group I, see the species election required below. If no additional fees are paid, Group I will be searched to the extent it reads on the first species of therapeutic agent named, a TNF antagonist.

Group II, claim(s) 24-35 and 37 in so far as they are drawn to IL-21/IL-21R agonist that is IL-21 polypeptide and method of treating or preventing a cancer or infectious disorder and method for increasing the ability of a vaccine composition containing an antigen to elicit a protective immune response in a subject against an antigen, and method of modulating the activity of an immune cell. If no additional fees are paid, Group III will be searched to the extent it reads on the first species of cytokine named, IL-21.

This application contains claims directed to more than one species of the generic invention. These species are deemed to lack unity of invention because they are not so linked as to form a single general inventive concept under PCT Rule 13.1.

In order for more than one species to be examined, the appropriate additional examination fees must be paid. The species are as follows:

For Group I, a species of therapeutic agent selected from the group consisting of: IL-12 antagonist, IL-15 antagonist, IL-17 antagonist, IL-18 antagonist, IL-22 antagonist, T cell depleting agent, B cell depleting agent, methotrexate, leflunomide, sirolimus (rapamycin) or analog thereof, Cox-2 inhibitor, a cPLA2 inhibitor, an NSAID, and a p38 inhibitor.

For Group II, species of cytokine selected from the group consisting of: GM-CSF and IL-15.

The claims are deemed to correspond to the species listed above in the following manner:

Therapeutic agent, claims 3, 4 and 49-54.

Cytokine, claim 33.

The following claim(s) are generic: 1, 2, 5-9, 13, 14, 24-27, 29, 34, 35, 37-48, 55-63.

The inventions listed as Groups I and II do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons: The inventions listed as Groups I-II do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons: Pursuant to 37 C.F.R. § 1.475(B-D), the IS/US considers that where multiple products and processes are claimed, the main invention shall consist of the first invention of the category first mentioned in the claims and the first recited invention of each of the other categories related thereto. Accordingly, the main invention (Group I) comprises the first recited product, IL-21/IL-21R antagonist that is extracellular domain of IL-21 receptor or fusion protein thereof and first use of the antagonist in a method of treatment. Group III does not share the same or corresponding technical feature because Group III is drawn to an agonist of IL-21R and different methods of treatment or use. This authority therefore considers that the three inventions do not share a special technical feature within the meaning of PCT Rule 13.2 and thus do not relate to a single general inventive concept within the meaning of PCT Rule 13.1.

The species listed above do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, the species lack the same or corresponding special technical features for the following reasons: they are different compounds having different structures and activities, and therefore do not share the same technical feature. This authority therefore considers that the several inventions do not share a special technical feature within the meaning of PCT Rule 13.2 and thus do not relate to a single general inventive concept within the meaning of PCT Rule 13.1.

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